

GenCore version 5.1.4.p5\_4578  
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OM nucleic - nucleic search, using sw model

Run on: April 23, 2003, 12:06:52 ; Search time 172.986 Seconds  
(without alignments)  
3059.318 Million cell updates/sec

Title: US-09-198-779b-1

Perfect score: 235  
Sequence: 1 gtttcgcgtcagctcgtcgt.....ctgcgaggtgtcaagcccc 235

Scoring table: IDENTITY-NUC

Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 50 summaries

Database :

N.Geneseq\_101002:\*

- 1: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1980.DAT:\*
- 2: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1981.DAT:\*
- 3: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1982.DAT:\*
- 4: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1983.DAT:\*
- 5: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1984.DAT:\*
- 6: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1985.DAT:\*
- 7: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1986.DAT:\*
- 8: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1987.DAT:\*
- 9: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1988.DAT:\*
- 10: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1989.DAT:\*
- 11: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1990.DAT:\*
- 12: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1991.DAT:\*
- 13: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1992.DAT:\*
- 14: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1993.DAT:\*
- 15: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1994.DAT:\*
- 16: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1995.DAT:\*
- 17: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1996.DAT:\*
- 18: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1997.DAT:\*
- 19: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1998.DAT:\*
- 20: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1999.DAT:\*
- 21: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2000.DAT:\*
- 22: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2001A.DAT:\*
- 23: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2001B.DAT:\*
- 24: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2002.DAT:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	121	51.5	2061	21	AAA48574
2	75.6	32.2	1674	21	AAC47744
3	65.8	28.0	1380	20	AAK07185
4	61.6	26.2	1182	19	AAT99143
5	61	26.0	1182	19	AAT99141
6	59.4	25.3	1182	19	AAT99142
7	54.6	23.2	399	20	AAK81128
8	53	22.6	1485	20	AAK07184
9	53	22.6	1518	21	AAA51037

10	51	21.7	1582	20	AAK07183	Corn S-adenosylmet
11	50.4	21.4	635	21	AAC44219	Arabidopsis thalia
12	50.4	21.4	1508	21	AAC33986	Arabidopsis thalia
13	50.4	21.4	1521	21	AAC45478	Arabidopsis thalia
14	50.4	21.4	1529	21	AAC33535	Arabidopsis thalia
15	49.8	21.2	1653	21	AAC45944	Arabidopsis thalia
16	49.8	21.2	1654	21	AAC35348	Arabidopsis thalia
17	46.6	19.8	1636	22	AAD02296	Nicotiana tabacum
18	42.4	18.0	1393	21	AAC46421	Arabidopsis thalia
19	42.4	18.0	1393	21	AAC33674	Arabidopsis thalia
20	40.2	17.1	1208	15	AA064204	snac gene encoding
21	40.2	17.1	5392	15	AA064201	Sequence comprisin
22	40	17.0	4848	24	AAD22684	Streptomyces fridi
23	40	17.0	4848	24	AAD22686	Streptomyces fridi
24	39.4	16.8	1693	21	AAK75637	Nucleotide sequenc
25	39.4	16.8	1693	21	AAK256005	Contig 004 from co
26	37.4	15.9	297	24	ABL72111	Corn tassell-derive
27	34.6	14.7	56485	21	AAK81476	N. meningitidis pa
28	34.6	14.7	349980	21	AAK21612	N. meningitidis pa
29	34.6	14.7	837096	21	AAK81489	N. meningitidis pa
30	34.4	14.6	4403765	22	AA199683	Mycobacterium tube
31	34.4	14.6	4411529	22	AA199682	Mycobacterium tube
32	33	14.0	17512	23	ABL09034	Drosophila melanog
33	31.2	13.3	1844	22	AAH99804	Human protein enco
34	30.8	13.1	66788	23	AAH59515	Protonibacterium
35	30.6	13.0	606	23	AAH51580	Pseudomonas aerugi
36	30.6	13.0	657	21	AAK12426	Aspergillus oryzae
37	30.6	13.0	2247	23	ABL29661	Drosophila melanog
38	30.2	12.9	566	21	AAK08527	Fusarium venenatum
39	29.8	12.7	1519	20	AAV64373	GABA-gated chlorid
40	29.6	12.6	3550	23	ABL09035	Drosophila melanog
41	29.4	12.5	1683	18	AAT86246	CDNA encoding mugw
42	29.2	12.4	1557	21	AAZ49567	Maize M109 protein
43	29.2	12.4	2604	21	AAA48576	CDNA encoding whea
44	29	12.3	1086	23	ABL04431	Drosophila melanog
45	29	12.3	1221	22	AAH66738	C glutaminc codin
46	29	12.3	1239	16	AAK04566	S-adenosylmethioni
47	29	12.3	1344	22	AAK71873	Corynebacterium gl
48	29	12.3	1344	23	AAK96132	C. glutamicum gene
49	29	12.3	3205	23	ABL04430	Drosophila melanog
50	29	12.3	349980	22	AAH68529	C glutaminc codin

#### ALIGNMENTS

RESULT 1  
AAA48574  
ID AAA48574 standard; CDNA: 2061 BP.

AAA48574;  
19-SEP-2000 (first entry)

CDNA encoding corn protein phosphatase 2A regulatory subunit A.  
Corn: protein phosphatase 2A; protein phosphorylation modulation;  
transgenic plant; gene therapy; ss.

Zea mays.

Key Location/Qualifiers  
CDS 56..1820  
/\*tag= a  
/\*product= "protein phosphatase 2A regulatory  
subunit A"

WO2000036121-A2.  
22-JUN-2000.  
15-DEC-1999; 99WO-US29823.

PR 16-DEC-1998; 98US-0112541.  
XX  
XX (DUPO ) DU PONT DE NEMOURS & CO E I.  
XX  
XX Famodu OO, Miao G, Sakai H, Lee J, Rafalski JA, Klein TW.  
PI WPI: 2000-431599/37.  
XX P-ESDB; AA199819.  
DR  
XX Polynucleotides encoding plant protein phosphatase useful for  
PT modulating reversible protein phosphorylation in plants -  
XX  
XX Claim 4; Page 53-54; 73pp; English.  
XX  
XX The present sequence encodes corn protein phosphatase 2A regulatory  
CC subunit A. The sequence was identified in clone p0018.chsug10r.fis of a  
CC cDNA library made from corn ear shoot. BLAST analysis showed that the  
CC present sequence encodes protein phosphatase 2A regulatory subunit A.  
CC The sequence may be used for the recombinant production of the protein  
CC in vivo, e.g. via a gene therapy protocol, or in vitro, e.g. in  
CC fermentation culture. The protein may then be used to modulate the  
CC process of reversible protein phosphorylation in plants. It may be used  
CC directly to supplement a plant's own production of the enzyme or to  
CC rectify mutations that result in the expression of inactive protein.  
CC The protein may also be used to test for modulators of protein  
CC phosphorylation which may be used to alter the activity of the enzyme.  
XX  
SQ Sequence 2061 BP; 549 A; 432 C; 498 G; 582 T; 0 other;  
  
Query Match 51.5%; Score 121; DB 21; Length 2061;  
Best Local Similarity 100.0%; Pred. No. 2.8e-31;  
Matches 121; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
OY 1 GTTTCCTGCTAGCCTGGTGCAGAGATCGACGCTGCCATCTATAATGAGCGTCC 60  
|||  
Db 1908 GTTTCCTGCTAGCCTGGTGCAGAGATCGACGCTGCCATCTATAATGAGCGTCC 1967  
|||  
OY 61 TGATCCATTGCTGCTGCTTTATTAATGATTAATGAGAGACCAACACGCTACT 120  
|||  
Db 1968 TGATCCATTGCTGCTGCTTTATTAATGATTAATGAGAGACCAACACGCTACT 2027  
|||  
OY 121 T 121  
||  
Db 2028 T 2028  
||  
  
RESULT 2  
AAC47744  
ID AAC47744 standard; DNA; 1674 BP.  
XX  
XX AAC47744;  
AC  
XX  
DT 18-OCT-2000 (first entry)  
XX  
XX Zea mays DNA fragment SEQ ID NO: 54954.  
DE  
XX  
XX Hybridisation assay; genetic mapping; gene expression control;  
KW protein identification; signal transduction pathway; metabolic;  
XX pathway; promoter; termination sequence; corn; ss.  
XX  
XX Zea mays subsp. mays.  
OS  
XX  
PN EPI033405-A2.  
XX  
XX  
PD 06-SEP-2000.  
XX  
PF 25-FEB-2000; 2000EP-0301439.  
XX  
PR 25-FEB-1999; 99US-0121825.  
PR 05-MAR-1999; 99US-0123180.  
PR 08-MAR-1999; 99US-0123548.  
PR 23-MAR-1999; 99US-0125788.  
PR 25-MAR-1999; 99US-0126264.

PR 29-MAR-1999; 99US-0126785.  
PR 01-APR-1999; 99US-0127462.  
PR 06-APR-1999; 99US-0128234.  
PR 08-APR-1999; 99US-0128714.  
PR 16-APR-1999; 99US-0129845.  
PR 19-APR-1999; 99US-0130077.  
PR 23-APR-1999; 99US-0130449.  
PR 23-APR-1999; 99US-0130510.  
PR 28-APR-1999; 99US-0130899.  
PR 30-APR-1999; 99US-0131449.  
PR 30-APR-1999; 99US-0132048.  
PR 04-APR-1999; 99US-0132407.  
PR 05-MAY-1999; 99US-0132485.  
PR 06-MAY-1999; 99US-0132486.  
PR 07-MAY-1999; 99US-0132487.  
PR 11-MAY-1999; 99US-0132863.  
PR 14-MAY-1999; 99US-0134256.  
PR 14-MAY-1999; 99US-0134218.  
PR 14-MAY-1999; 99US-0134219.  
PR 14-MAY-1999; 99US-0134221.  
PR 14-MAY-1999; 99US-0134370.  
PR 18-MAY-1999; 99US-0134376.  
PR 19-MAY-1999; 99US-0134941.  
PR 20-MAY-1999; 99US-0135124.  
PR 21-MAY-1999; 99US-0135124.  
PR 24-MAY-1999; 99US-0135353.  
PR 25-MAY-1999; 99US-0135629.  
PR 27-MAY-1999; 99US-0136021.  
PR 28-MAY-1999; 99US-0136392.  
PR 01-JUN-1999; 99US-0136782.  
PR 03-JUN-1999; 99US-0137222.  
PR 04-JUN-1999; 99US-0137528.  
PR 07-JUN-1999; 99US-0137502.  
PR 08-JUN-1999; 99US-0137724.  
PR 10-JUN-1999; 99US-0138094.  
PR 10-JUN-1999; 99US-0138540.  
PR 14-JUN-1999; 99US-0139116.  
PR 16-JUN-1999; 99US-0139452.  
PR 16-JUN-1999; 99US-0139453.  
PR 17-JUN-1999; 99US-0139492.  
PR 18-JUN-1999; 99US-0139454.  
PR 18-JUN-1999; 99US-0139455.  
PR 18-JUN-1999; 99US-0139456.  
PR 18-JUN-1999; 99US-0139457.  
PR 18-JUN-1999; 99US-0139458.  
PR 18-JUN-1999; 99US-0139459.  
PR 18-JUN-1999; 99US-0139460.  
PR 18-JUN-1999; 99US-0139461.  
PR 18-JUN-1999; 99US-0139462.  
PR 18-JUN-1999; 99US-0139463.  
PR 18-JUN-1999; 99US-0139750.  
PR 18-JUN-1999; 99US-0139763.  
PR 21-JUN-1999; 99US-0139817.  
PR 22-JUN-1999; 99US-0139899.  
PR 23-JUN-1999; 99US-0140353.  
PR 23-JUN-1999; 99US-0140354.  
PR 24-JUN-1999; 99US-0140699.  
PR 28-JUN-1999; 99US-0140823.  
PR 29-JUN-1999; 99US-0140991.  
PR 30-JUN-1999; 99US-0141287.  
PR 01-JUL-1999; 99US-0141842.  
PR 01-JUL-1999; 99US-0142154.  
PR 02-JUL-1999; 99US-0142055.  
PR 06-JUL-1999; 99US-0142390.  
PR 08-JUL-1999; 99US-0142803.  
PR 09-JUL-1999; 99US-0142920.  
PR 12-JUL-1999; 99US-0142977.  
PR 13-JUL-1999; 99US-0143624.  
PR 14-JUL-1999; 99US-0143624.  
PR 15-JUL-1999; 99US-0144005.  
PR 16-JUL-1999; 99US-0144085.  
PR 16-JUL-1999; 99US-0144086.



```

XX  Abell LM, Allen SM, Falco SC, Hitz WD, Kinney AJ;
PI  Rafalski JA, Thorpe CJ;
XX
DR  WPI: 1999-070263/06.
DR  P-PSDB: AAM97744.
XX
PT  New plant amino acid biosynthetic enzymes, DNA and chimeric genes -
PT  encode: dihydropicolinate reductase; diaminopimelate epimerase;
PT  threonine synthase; threonine deaminase; S-adenosylmethionine
PT  synthetase
XX
PS  Claim 47; Page 69-70; 98pp; English.
XX
CC  This is the nucleotide sequence of a cDNA contig that codes for
CC  a full-length wheat S-adenosylmethionine synthetase (see AAM97744).
CC  The contig was assembled from clones isolated from wheat kernel,
CC  leaf, seedling and root cDNA libraries and identified by comparison
CC  to public sequence databases using BLAST algorithms. It shows
CC  sequence similarity to the barley enzyme. The invention relates to
CC  new isolated nucleic acid fragments (see AAX07168-85) encoding plant
CC  enzymes (see AAM97727-44) that catalyze steps in the biosynthesis of
CC  lysine, threonine, methionine, cysteine and isoleucine from
CC  aspartate, the enzyme being selected from dihydropicolinate reductase,
CC  diaminopimelate epimerase, threonine synthase, threonine deaminase
CC  or S-adenosylmethionine synthetase. The invention also relates to
CC  the construction of a chimeric gene encoding all or a portion of
CC  the biosynthetic pathway enzyme, in sense or antisense orientation,
CC  where expression of the chimeric gene results in production of
CC  altered levels of the enzyme in a transformed host cell.
CC  Overexpression or reduction of expression of genes encoding the
CC  amino acid biosynthetic pathway enzymes in crop plants such as
CC  corn, soybean and wheat can be used to alter levels of the amino
CC  acids in human food and animal feed. Transformed host cells can
CC  also be used to identify compounds that inhibit one of the enzymes.
XX
SQ  Sequence 1380 BP; 299 A; 430 C; 379 G; 267 T; 5 other;

Query Match      28.0%; Score 65.8; DB 20; Length 1380;
Best Local Similarity 90.9%; Pred. No. 2.4e-12;
Matches 70; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

OY  159 CAGGTCATCAAGACCGCGCATAGCGCACTTGGCCGTAGCGAGCGGCACTTCACTG 218
    |||||||
DB  1155 CAGGTCATCAAGACCGCGCTTACGTCCTTGGCCGTAGCGGCACTTCACTG 1214
OY  219 CGAGTGCTCAAGCCCC 235
    |||||||
DB  1215 GGAGTGCTGAAGCCCC 1231

RESULT 4
AAT99143
ID  AAT99143 standard; cDNA to mRNA; 1182 BP.
XX
AC  AAT99143;
XX
DT  26-MAR-1998 (first entry)
XX
DE  S-adenosylmethionine synthase 3 gene.
XX
KW  S-adenosylmethionine synthase 3 gene; barley; alkali resistant plant;
KW  sam3; ss.
XX
OS  Hordeum vulgare.
XX
PN  JP09313186-A.
XX
PD  09-DEC-1997.
XX
PE  28-MAY-1996; 96JP-0133406.
XX
PR  28-MAY-1996; 96JP-0133406.

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XX  (NIOC ) NIPPON OIL CO LTD.
XX
DR  WPI: 1998-080077/08.
DR  P-PSDB: AAM34542.
XX
PT  S-adenosyl-methionine synthase gene - useful in producing plants
PT  resistant to alkaline soil
XX
PS  Claim 6; Page 10-11; 13pp; Japanese.
XX
CC  This sequence represents the S-adenosylmethionine synthase 3 (sam3)
CC  gene. This DNA sequence may be used in a vector to produce plants which
CC  are resistant to alkaline soil.
XX
SQ  Sequence 1182 BP; 253 A; 368 C; 331 G; 230 T; 0 other;

Query Match      26.2%; Score 61.6; DB 19; Length 1182;
Best Local Similarity 88.2%; Pred. No. 6.3e-11;
Matches 67; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

OY  160 AGGTCATCAAGACCGCGCATAGCGCACTTGGCCGTAGCGAGCGGCACTTCACTGC 219
    |||||||
DB  1084 AGGTCATCAAGACCGCGCTTATAGTCATCTTGGCCGCGAGATGCCACTTCACTGG 1143
OY  220 GAGTGCTCAAGCCCC 235
    |||||||
DB  1144 GAGTGCTGAAGCCCC 1159

RESULT 5
AAT99141
ID  AAT99141 standard; cDNA to mRNA; 1182 BP.
XX
AC  AAT99141;
XX
DT  26-MAR-1998 (first entry)
XX
DE  S-adenosylmethionine synthase 1 gene.
XX
KW  S-adenosylmethionine synthase 1 gene; barley; alkali resistant plant;
KW  sam1; ss.
XX
OS  Hordeum vulgare.
XX
PN  JP09313186-A.
XX
PD  09-DEC-1997.
XX
PE  28-MAY-1996; 96JP-0133406.
XX
PR  28-MAY-1996; 96JP-0133406.
XX
PA  (NIOC ) NIPPON OIL CO LTD.
XX
DR  WPI: 1998-080077/08.
DR  P-PSDB: AAM34540.
XX
DE  S-adenosyl-methionine synthase gene - useful in producing plants
DE  resistant to alkaline soil
XX
PS  Claim 4; Page 8-9; 13pp; Japanese.
XX
CC  This sequence represents the S-adenosylmethionine synthase 1 (sam1)
CC  gene. This DNA sequence may be used in a vector to produce plants which
CC  are resistant to alkaline soil.
XX
SQ  Sequence 1182 BP; 253 A; 372 C; 324 G; 233 T; 0 other;

Query Match      26.0%; Score 61; DB 19; Length 1182;
Best Local Similarity 87.0%; Pred. No. 1e-10;
Matches 67; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

```



QY 159 CAGGTTTCATCAGACCGCCGATACGGCCATTTGGCCGTGACGAGCCGCACTTCACCTG 218  
 |||||||  
 DB 1083 CAGGTTTCATCAGACGCTGCTTACGCTCATTGGCCGCGATGATGCTGACTTCACCTG 1142  
 |||||||  
 QY 219 CGAGGTGCTCAAGCCCC 235  
 |||||||  
 DB 1143 GGAGGTGCTGAAGCCCC 1159

## RESULT 6

AAAT99142  
 ID AAAT99142 standard; cDNA to mRNA; 1182 BP.

AAAT99142;

26-MAR-1998 (first entry)

S-adenosylmethionine synthase 2 gene.

S-adenosylmethionine synthase 2 gene; barley; alkali resistant plant;

sam2; ss.

Hordeum vulgare.

JP09313186-A.

09-DEC-1997.

28-MAY-1996; 96JP-0133406.

28-MAY-1996; 96JP-0133406.

(NIOC ) NIPPON OIL CO LTD.

WPI: 1998-080077/08.

P-PSDB; AAM34541.

S-adenosyl-methionine synthase gene - useful in producing plants  
 resistant to alkaline soil

Claim 5; Page 9-10; 13pp; Japanese.

This sequence represents the S-adenosylmethionine synthase 2 (sam2)  
 CC gene. This DNA sequence may be used in a vector to produce plants which  
 are resistant to alkaline soil.

Sequence 1182 BP; 261 A; 368 C; 324 G; 229 T; 0 other;

Query Match 25.3%; Score 59.4; DB 19; Length 1182;

Best Local Similarity 85.7%; Pred. No. 3,6e-10;

Matches 66; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

QY 159 CAGGTTTCATCAGACCGCCGATACGGCCATTTGGCCGTGACGAGCCGCACTTCACCTG 218  
 |||||||  
 DB 1083 CAGGTTTCATCAGACGCTGCTTACGCTCATTGGCCGCGATGATGCTGACTTCACCTG 1142  
 |||||||

QY 219 CGAGGTGCTCAAGCCCC 235  
 |||||||

DB 1143 GGAGGTGCTGAAGCCCC 1159

## RESULT 7

AAAX81128  
 ID AAAX81128 standard; cDNA; 399 BP.

AAAX81128;

06-SEP-1999 (first entry)

S-adenosyl methionine (SAM) partial cDNA sequence.

S-adenosyl methionine (SAM) partial cDNA sequence. sark; sam gene; sag;  
 senescence-associated gene; plant senescence; promoter; pharmaceutical;

KW plant maturation; S-adenosyl methionine; flower; fruit development; ss.  
 XX Phaseolus vulgaris.  
 OS

PN WO9929159-A1.

17-JUN-1999.

08-DEC-1998; 98WO-US25799.

08-DEC-1997; 97US-0067898.

(VITA-) VITALITY BIOTECHNOLOGIES INC.

Gepstein S, Hajuoje T, Rosner A;

WPI: 1999-404873/34.

P-PSDB: AAY21978.

DNA encoding senescence-associated genes for a senescence  
 receptor-like protein kinase

Claim 18; Fig 4; 70pp; English.

The invention relates to a senescence-associated receptor-like protein  
 kinase (sark) gene. The sark gene is a senescence-associated gene (sag)  
 and is expressed early in the plant senescence process. The sark gene  
 promoter is useful for driving expression of foreign genes having a  
 desired product, such as a pharmaceutical, during the process of plant  
 maturation. The sark gene promoter can be used to drive expression of  
 resistance genes against pathogens or pests during senescence when the  
 plant is particularly susceptible to infection or infestation. The sark  
 gene promoter may also be used to drive expression of a gene encoding an  
 inhibitor of senescence. Plant senescence may be inhibited by use of  
 antisense sark constructs. Over expression of the sag genes, using the  
 sark or sam (S-adenosyl methionine) gene promoters is useful for  
 induction of early senescence. This is useful to obtain flower or fruit  
 development prior to specific pest onset, prior to undesirable cross-  
 fertilization from related crops, at a specific time during storage or  
 retail, or to avoid development of plant structures that are not of  
 agronomic importance. The present sequence represents a partial cDNA  
 clone of S-adenosyl methionine (SAM).

Sequence 399 BP; 108 A; 80 C; 82 G; 129 T; 0 other;

Query Match 23.2%; Score 54.6; DB 20; Length 399;

Best Local Similarity 81.8%; Pred. No. 1e-08;

Matches 63; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 159 CAGGTTTCATCAGACCGCCGATACGGCCATTTGGCCGTGACGAGCCGCACTTCACCTG 218  
 |||||||  
 DB 101 CAGGTTCTGAAGACGCTGATATGACACTTCGCGACAGAGAGCGCTGACTTCACATG 160  
 |||||||

QY 219 CGAGGTGCTCAAGCCCC 235  
 |||||||

DB 161 GGAAGTGTCTCAAGCCCC 177

## RESULT 8

AAAX07184  
 ID AAAX07184 standard; cDNA; 1485 BP.

AAAX07184;

21-MAY-1999 (first entry)

Soybean S-adenosylmethionine synthetase cDNA clone s2.12D06.

S-adenosylmethionine synthetase; soybean; amino acid; lysine;  
 threonine; methionine; cysteine; isoleucine; transgenic plant;  
 crop improvement; food; feedstuff; ss.

Glycine max.

```

XX Key Location/Qualifiers
FH CDS 74..1252
FT /*tag= a
XX
XX PN W09855601-A2.
XX
XX PD 10-DEC-1998.
XX
XX PF 05-JUN-1998; 98WO-US11692.
XX
XX PR 12-JUN-1997; 97US-0049443.
XX PR 06-JUN-1997; 97US-0048771.
XX
XX PA (DUPO ) DU PONT DE NEMOURS & CO E I.
XX
XX PI Abell LM, Allen SM, Falco SC, Hiltz WD, Kinney AJ;
XX PI Rafalski JA, Thorpe CJ;
XX
XX DR WPI, 1999-070263/06.
XX DR P-PSDB; AAM97743.
XX
XX PS New plant amino acid biosynthetic enzymes, DNA and chimeric genes -
XX PT encode: dihydropicolinate reductase; diaminopimelate epimerase;
XX PT threonine synthase; threonine deaminase; S-adenosylmethionine
XX PT synthetase
XX
XX PS Claim 44; Page 66-67; 98pp; English.
XX
XX CC This is the nucleotide sequence of cDNA clone 52.12b06, which
XX CC codes for a full-length soybean S-adenosylmethionine synthetase
XX CC (see AAM97743). The clone was isolated from a soybean seed cDNA
XX CC library and identified by comparison to public sequence databases
XX CC using BLAST algorithms. It shows sequence similarity to the
XX CC tomato enzyme. The invention relates to new isolated nucleic
XX CC acid fragments (see AAX07168-85) encoding plant enzymes (see
XX CC AAM97727-44) that catalyze steps in the biosynthesis of lysine,
XX CC threonine, methionine, cysteine and isoleucine from aspartate, the
XX CC enzyme being selected from dihydropicolinate reductase,
XX CC diaminopimelate epimerase, threonine synthase, threonine deaminase
XX CC or S-adenosylmethionine synthetase. The invention also relates to
XX CC the construction of a chimeric gene encoding all or a portion of
XX CC the biosynthetic pathway enzyme, in sense or antisense orientation,
XX CC where expression of the chimeric gene results in production of
XX CC altered levels of the enzyme in a transformed host cell.
XX CC Overexpression or reduction of expression of genes encoding the
XX CC amino acid biosynthetic pathway enzymes in crop plants such as
XX CC corn, soybean and wheat can be used to alter levels of the amino
XX CC acids in human food and animal feed. Transformed host cells can
XX CC also be used to identify compounds that inhibit one of the enzymes.
XX
XX SQ Sequence 1485 BP; 366 A; 373 C; 357 G; 389 T; 0 other;
XX
XX Query Match 22.6%; Score 53; DB 20; Length 1485;
XX Best Local Similarity 80.5%; Pred. No. 6.4e-08;
XX Matches 62; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

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XX XX Soybean S-adenosyl-L-methionine synthetase cDNA.
DE S-adenosyl-L-methionine synthetase; SAMS; probe; promoter; embryo;
XX constitutive; tissue-specific; development-specific;
XX herbicide resistance; pathogen resistance; ss.
XX
XX OS Glycine max.
XX
XX FH Key Location/Qualifiers
XX FT 5'UTR 1..73
XX FT /*tag= a
XX FT CDS 74..1252
XX FT /*tag= b
XX FT /product= S-adenosyl-L-methionine_synthetase
XX
XX PN W0200037662-A2.
XX
XX PD 29-JUN-2000.
XX
XX PF 17-DEC-1999; 99WO-US30180.
XX
XX PR 21-DEC-1998; 98US-0113045.
XX
XX PA (DUPO ) DU PONT DE NEMOURS & CO E I.
XX
XX PI Falco SC, Li Z;
XX PI WPI, 2000-442682/38.
XX
XX DR S-adenosyl-L-methionine synthetase promoter for expressing target
XX PT heterologous herbicide-resistance or pathogen-resistance nucleic acid
XX PT fragments in plants, especially soybean
XX
XX PS Example 2; Page 39; 50pp; English.
XX
XX CC This is the soybean full-length S-adenosyl-L-methionine synthetase (SAMS)
XX CC cDNA, which was used to generate a probe to isolate a SAMS promoter. The
XX CC SAMS promoter is active in seedlings and callus and over-expression of a
XX CC gene in embryo stage can be achieved at an early developing stage using
XX CC the SAMS promoter. The SAMS promoter may be used as an alternative to
XX CC cauliflower mosaic virus 35S promoter to drive expression of selectable
XX CC marker genes. Plant cells transformed with the SAMS constitutive promoter
XX CC are useful for increasing or decreasing the expression of heterologous
XX CC nucleic acid fragments in a plant, preferably corn, rice, wheat, barley,
XX CC palm, Arabidopsis, soybean, oil seed Brassica, peanut, sunflower,
XX CC safflower, cotton, tobacco, tomato, potato or cocoa. Target heterologous
XX CC nucleic acid fragments include herbicide or pathogen resistance
XX CC nucleic acid fragments.
XX
XX SQ Sequence 1518 BP; 399 A; 373 C; 357 G; 389 T; 0 other;
XX
XX Query Match 22.6%; Score 53; DB 21; Length 1518;
XX Best Local Similarity 80.5%; Pred. No. 6.5e-08;
XX Matches 62; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

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RESULT 9
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XX AC AAX51037;
XX
XX DT 09-OCT-2000 (first entry)

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RESULT 10
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ID AAX07183 standard; cDNA: 1582 BP.
XX
XX AC AAX07183;
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XX DT 21-MAY-1999 (first entry)

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XX Hybridisation assay; genetic mapping; gene expression control;  
KW protein identification; signal transduction pathway;  
KW metabolic pathway; promoter; termination sequence; ss.  
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OS Arabidopsis thaliana.  
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KW	Hybridisation assay; genetic mapping; gene expression control	
XX		
KM	protein identification; signal transduction pathway;	
XX	metabolic pathway; promoter; termination sequence; ss.	
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XX		
PN	EP1033405-A2.	
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PR 29-OCT-1999; 99US-0162142.

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Query Match 21.2%; Score 49.8; DB 21; Length 1654;
Best Local Similarity 77.9%; Pred. No. 8.5e-07;
Matches 60; Conservative 0; Mismatches 17; Indels 0; Gaps 0;

QY 159 CAGGTTTCATCAAGACCGCGCATACGCCACTTGGCCGTGACGACGCCGACTTCACCTG 218
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Db 1228 CAGGTTCCAGAAACCGCTGCGTATGCGCATTTCCGCGCTATGACCTGACTTCACCTTG 1287

QY 219 CGAGGTGTCAGACCCC 235
    |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  ||
Db 1288 GGAGGTGTCTCAAGCCGC 1304

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RESULT 17
AAD02296
ID AAD02296 standard; DNA; 1636 BP.
XX
AC AAD02296;
XX
DT 28-MAR-2001 (first entry)
XX
DE Nicotiana tabacum S-adenosylmethionine synthetase (SAMS) DNA.
XX
KM Tobacco: alkaloid; nicotine; transgenic plant; pharmaceutical protein;
KW herbicide resistance; S-adenosylmethionine synthetase; SAMS; ds.
XX
OS Nicotiana tabacum.
XX
FT key Location/Qualifiers
FT CDS 96..1268
FT /*tag= a

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FT FT /product= "Nicotiana tabacum S-adenosylmethionine
FT FT synthetase (SAMS) protein"
XX
XX WO200067558-A1.
XX
XX 16-NOV-2000.
XX
XX 05-MAY-2000; 2000WO-US12450.
XX
XX 06-MAY-1999; 99US-0132919.
XX
XX (TIMK/) TIMKO M.
XX
XX Timko M;
XX
XX WPI: 2001-007279/01.
XX
XX P-PSDB: AAV72078.
XX
XX New nucleic acid encoding alkaloid-synthesis enzymes in tobacco, useful
XX e.g. for producing transgenic plants with altered nicotine content
XX
XX Claim 1; Page 78-79; 103pp; English.
XX
XX The invention relates to enzymes involved in alkaloid, specifically
XX nicotine, synthesis in tobacco and nucleic acids encoding them. The
XX nucleic acid of the invention can be used, in sense or antisense
XX orientation, to produce transgenic tobacco plants with altered
XX alkaloid content, and also for expression of exogenous proteins,
XX e.g. pharmaceutical proteins or proteins implicated in resistance
XX to herbicides. The protein of the invention can be used to
XX identify modulators of enzymatic activity in plants.
XX The present sequence is Nicotiana tabacum S-adenosylmethionine
XX synthetase (SAMS) DNA. This enzyme is involved in the nicotine
XX biosynthetic pathway.
XX
XX Sequence 1636 BP; 444 A; 365 C; 373 G; 454 T; 0 other;

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Query Match 19.8%; Score 46.6; DB 22; Length 1636;
Best Local Similarity 75.3%; Pred. No. 1.1e-05;
Matches 58; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

QY 159 CAGGTTTCATCAAGACCGCGCATACGCCACTTGGCCGTGACGACGCCGACTTCACCTG 218
    |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  ||
Db 1175 CAGGTACCAAGAGACTGAGCTTATGTCACCTTGGCCGTGATGACCCGACTTCATG 1234

QY 219 CGAGGTGTCAGACCCC 235
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Db 1235 GGAGACTGTCAAGCTCC 1251

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RESULT 18
AAC46421
ID AAC46421 standard; DNA; 1393 BP.
XX
AC AAC46421;
XX
XX 18-OCT-2000 (first entry)
XX
XX Arabidopsis thaliana DNA fragment SEQ ID NO: 50076.
XX
XX Hybridisation assay; genetic mapping; gene expression control;
XX protein identification; signal transduction pathway;
XX metabolic pathway; promoter; termination sequence; ss.
XX
XX Arabidopsis thaliana.
XX
XX EP1033405-A2.
XX
XX 06-SEP-2000.
XX
XX 25-FEB-2000; 2000EP-0301439.
XX
XX 25-FEB-1999; 99US-0121825.

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PR 05-MAR-1999; 99US-0121180.  
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PR 25-MAR-1999; 99US-0126264.  
PR 29-MAR-1999; 99US-0126785.  
PR 01-APR-1999; 99US-0127462.  
PR 06-APR-1999; 99US-0128234.  
PR 08-APR-1999; 99US-0128714.  
PR 16-APR-1999; 99US-0129845.  
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PR 21-APR-1999; 99US-0130449.  
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PR 23-APR-1999; 99US-0130891.  
PR 28-APR-1999; 99US-0131449.  
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PR 01-JUL-1999; 99US-0141842.  
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PR 13-JUL-1999; 99US-0143542.

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Query Match 18.0%; Score 42.4; DB 21; Length 1393;
Best Local Similarity 72.4%; Pred. No. 0.00028;
Matches 55; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

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Db 1175 AGGTTTCAGAAACGCGCATGTGACATTTGCGAAGAGACGACCTTCACCTGC 1234
    |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

OY 220 GAGGTGTCAGAGCCGC 235
    |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
Db 1235 GAGGTGTCAGAGCCGC 1250
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RESULT 19
AAC33674
ID AAC33674 standard; DNA: 1395 BP.
XX
AC AAC33674;
XX
DT 17-OCT-2000 (first entry)
XX
DE Arabidopsis thaliana DNA fragment SEQ ID NO: 3906.
XX
KW Hybridisation assay; genetic mapping; gene expression control;
KW protein identification; signal transduction pathway;
KW metabolic pathway; promoter; termination sequence; ss.
XX
OS Arabidopsis thaliana.
XX
XX EPI033405-A2.
XX
PD 06-SEP-2000.
XX
PF 25-FEB-2000; 2000EP-0301439.
XX
PR 25-FEB-1999; 9905-0121825.
PR 05-MAR-1999; 9905-0123180.
PR 09-MAR-1999; 9905-0123548.
PR 23-MAR-1999; 9905-0125788.
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PR 16-APR-1999; 9905-0129845.
PR 19-APR-1999; 9905-0130077.
PR 21-APR-1999; 9905-0130449.
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PR 06-MAY-1999; 9905-0132487.
PR 07-MAY-1999; 9905-0132863.
PR 11-MAY-1999; 9905-0134256.
PR 14-MAY-1999; 9905-0134218.
PR 14-MAY-1999; 9905-0134219.
PR 14-MAY-1999; 9905-0134221.
PR 14-MAY-1999; 9905-0134370.
PR 18-MAY-1999; 9905-0134768.
PR 19-MAY-1999; 9905-0134941.
PR 20-MAY-1999; 9905-0135124.
PR 21-MAY-1999; 9905-0135353.
PR 24-MAY-1999; 9905-0135629.
PR 25-MAY-1999; 9905-0136021.
PR 27-MAY-1999; 9905-0136392.
PR 28-MAY-1999; 9905-0136782.
PR 01-JUN-1999; 9905-0137222.
PR 03-JUN-1999; 9905-0137528.
PR 04-JUN-1999; 9905-0137502.
PR 07-JUN-1999; 9905-0137724.
PR 08-JUN-1999; 9905-0138094.
PR 10-JUN-1999; 9905-0138540.
PR 10-JUN-1999; 9905-0138847.
PR 14-JUN-1999; 9905-0139119.
PR 16-JUN-1999; 9905-0139452.
PR 16-JUN-1999; 9905-0139453.
PR 17-JUN-1999; 9905-0139454.
PR 18-JUN-1999; 9905-0139454.
PR 18-JUN-1999; 9905-0139455.
PR 18-JUN-1999; 9905-0139456.
PR 18-JUN-1999; 9905-0139457.
PR 18-JUN-1999; 9905-0139458.
PR 18-JUN-1999; 9905-0139459.
PR 18-JUN-1999; 9905-0139460.
PR 18-JUN-1999; 9905-0139461.
PR 18-JUN-1999; 9905-0139462.
PR 18-JUN-1999; 9905-0139463.
PR 18-JUN-1999; 9905-0139750.
PR 18-JUN-1999; 9905-0139763.
PR 21-JUN-1999; 9905-0139817.
PR 22-JUN-1999; 9905-0139859.
PR 23-JUN-1999; 9905-0140353.
PR 23-JUN-1999; 9905-0140354.
PR 24-JUN-1999; 9905-0140695.
PR 28-JUN-1999; 9905-0140823.
PR 29-JUN-1999; 9905-0140991.
PR 30-JUN-1999; 9905-0141287.
PR 01-JUL-1999; 9905-0141842.
PR 01-JUL-1999; 9905-0142154.
PR 02-JUL-1999; 9905-0142055.
PR 06-JUL-1999; 9905-0142390.
PR 06-JUL-1999; 9905-0142803.
PR 09-JUL-1999; 9905-0142920.
PR 12-JUL-1999; 9905-0142977.
PR 13-JUL-1999; 9905-0143542.
PR 14-JUL-1999; 9905-0143620.
PR 15-JUL-1999; 9905-0144005.
PR 16-JUL-1999; 9905-0144085.
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PR 16-JUL-1999; 99US-0144086.
PR 19-JUL-1999; 99US-0144325.
PR 19-JUL-1999; 99US-0144331.
PR 19-JUL-1999; 99US-0144332.
PR 19-JUL-1999; 99US-0144333.
PR 19-JUL-1999; 99US-0144334.
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PR 20-JUL-1999; 99US-0144332.
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PR 21-JUL-1999; 99US-0145087.
PR 22-JUL-1999; 99US-0145085.
PR 22-JUL-1999; 99US-0145087.
PR 22-JUL-1999; 99US-0145089.
PR 22-JUL-1999; 99US-0145192.
PR 23-JUL-1999; 99US-0145145.
PR 23-JUL-1999; 99US-0145218.
PR 23-JUL-1999; 99US-0145224.
PR 26-JUL-1999; 99US-0145276.
PR 27-JUL-1999; 99US-01452913.
PR 27-JUL-1999; 99US-0145918.
PR 27-JUL-1999; 99US-0145919.
PR 28-JUL-1999; 99US-0145951.
PR 02-AUG-1999; 99US-0146386.
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PR 05-AUG-1999; 99US-0147260.
PR 06-AUG-1999; 99US-0147303.
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PR 09-AUG-1999; 99US-0147493.
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PR 10-AUG-1999; 99US-0148171.
PR 11-AUG-1999; 99US-0148319.
PR 12-AUG-1999; 99US-0148341.
PR 13-AUG-1999; 99US-0148565.
PR 13-AUG-1999; 99US-0148684.
PR 16-AUG-1999; 99US-0149368.
PR 17-AUG-1999; 99US-0149175.
PR 18-AUG-1999; 99US-0149426.
PR 20-AUG-1999; 99US-0149722.
PR 20-AUG-1999; 99US-0149723.
PR 20-AUG-1999; 99US-0149929.
PR 23-AUG-1999; 99US-0149902.
PR 23-AUG-1999; 99US-0149930.
PR 25-AUG-1999; 99US-0150566.
PR 26-AUG-1999; 99US-0150884.
PR 27-AUG-1999; 99US-0151065.
PR 27-AUG-1999; 99US-0151066.
PR 27-AUG-1999; 99US-0151080.
PR 30-AUG-1999; 99US-0151303.
PR 31-AUG-1999; 99US-0151438.
PR 01-SEP-1999; 99US-0151930.
PR 07-SEP-1999; 99US-0152363.
PR 10-SEP-1999; 99US-0153070.
PR 13-SEP-1999; 99US-0153758.
PR 15-SEP-1999; 99US-0154039.
PR 16-SEP-1999; 99US-0154779.
PR 20-SEP-1999; 99US-0155139.
PR 22-SEP-1999; 99US-0155139.
PR 23-SEP-1999; 99US-0155486.
PR 24-SEP-1999; 99US-0155659.
PR 28-SEP-1999; 99US-0156458.
PR 29-SEP-1999; 99US-0156596.
PR 04-OCT-1999; 99US-0157117.
PR 05-OCT-1999; 99US-0157753.
PR 06-OCT-1999; 99US-0157865.
PR 07-OCT-1999; 99US-0158029.

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PR 08-OCT-1999; 99US-0158232.
PR 12-OCT-1999; 99US-0158369.
PR 13-OCT-1999; 99US-0159293.
PR 13-OCT-1999; 99US-0159294.
PR 13-OCT-1999; 99US-0159294.
PR 13-OCT-1999; 99US-0159295.
PR 14-OCT-1999; 99US-0159329.
PR 14-OCT-1999; 99US-0159330.
PR 14-OCT-1999; 99US-0159331.
PR 14-OCT-1999; 99US-0159637.
PR 14-OCT-1999; 99US-0159637.
PR 18-OCT-1999; 99US-0159638.
PR 18-OCT-1999; 99US-0159584.
PR 21-OCT-1999; 99US-0160741.
PR 21-OCT-1999; 99US-0160767.
PR 21-OCT-1999; 99US-0160768.
PR 21-OCT-1999; 99US-0160770.
PR 21-OCT-1999; 99US-0160814.
PR 21-OCT-1999; 99US-0160814.
PR 22-OCT-1999; 99US-0160819.
PR 22-OCT-1999; 99US-0160980.
PR 22-OCT-1999; 99US-0160981.
PR 22-OCT-1999; 99US-0160981.
PR 25-OCT-1999; 99US-0161404.
PR 25-OCT-1999; 99US-0161405.
PR 25-OCT-1999; 99US-0161405.
PR 25-OCT-1999; 99US-0161406.
PR 26-OCT-1999; 99US-0161359.
PR 26-OCT-1999; 99US-0161360.
PR 26-OCT-1999; 99US-0161361.
PR 28-OCT-1999; 99US-0161920.
PR 28-OCT-1999; 99US-0161992.
PR 28-OCT-1999; 99US-0161993.
PR 29-OCT-1999; 99US-0162142.

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Query Match      18.0%; Score 42.4; DB 21; Length 1395;
Best Local Similarity 72.4%; Pred. No. 0.00028;
Matches 55; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

```

```

QY 160 AGCTTCATCAGACCGCGCATACGCCACTTGGCCCGTGACGACGCGGACTTCACTTC 219
Db 1175 AGCTTTCAGAAAACGCGACGCGTATGACATTTGGAGAGACGACCGCTGACTTCACTTCG 1234
QY 220 GAGGTGCTGACGCCCC 235
Db 1235 GAGGTGCTGACGCCAC 1250

```

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RESULT 20
AA064204
ID AA064204 standard; cDNA; 1208 BP.
XX
AC AA064204;
XX
DT 18-NOV-1994 (first entry)
XX
DE snac gene encoding enzyme in streptogramin biosynthetic pathway.
XX
KW Antibiotic; streptogramin; snac; snab; snac; biosynthesis; enzyme;
KM biosynthetic pathway; Streptomyces pristinaespiralis; ds.
XX
OS Streptomyces pristinaespiralis.
XX
FH Key Location/Qualifiers
FT CDS 1..1209
FT FT /*tag= a
XX
PN FR2696189-A.
PD 01-APR-1994.
XX
PF 25-SEP-1992; 92FR-0011441.
XX
XX 25-SEP-1992; 92FR-0011441.
XX
XX (RHON ) RHONE POULENC RORER SA.
XX

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[illegible]

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PS prodn. or bio-conversion of streptogramin(s) or prodn. of
PT streptogramin intermediates, derivs. or hybrid antibiotics
XX
XX Disclosure; page 44-47; 83pp; French.
CC This sequence comprises the snaA, snab and snac genes which are
CC involved in the biosynthesis of streptogramins, antibiotics active
CC against Gram-positive bacteria. The identification of the sequences
CC encoding the enzymes involved in the biosynthetic pathway means that
CC they can be isolated and manipulated. Mutant microorganisms in
CC which a step in the streptogramin biosynthetic pathway is blocked
CC can be cultured to produce streptogramin intermediates, which may
CC later be converted to streptogramin derivatives. Recombinant cells
CC may also be used for the bioconversion of streptogramins from one
CC form to another or for the production of hybrid antibiotics.
SQ Sequence 5392 BP; 811 A; 2161 C; 1671 G; 749 T; 0 other:
Query Match      17.1%; Score 40..2; DB 15; Length 5392;
Best Local Similarity 46.7%; Pred. No. 0.0029;
Matches 57; Conservative 0; Mismatches 65; Indels 0; Gaps 0;
QY   113 ACCTACGTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNAGGTTCATCAGA 172
Db    4612 ACCGCGCCCCGGCCCATCATCGGCACCCTGCACTGCTGGCGCATCTACGCCGCCA 4671
QY   173 CC GCCGATTACGGCCACTTGGCCGCTGACGACGCCGACTTCACCTGCAGGTGCTCAAG 232
Db    4672 CCGCGGCTACGCCACTTGCGCGCGAACGACCCGACTTCACCTGGAGAGCGACCGACC 4731
QY       233 CC 234
Db    4732 GC 4733
RESULT 22
AAD22684 ID AAD22684 standard; DNA: 4848 BP.
XX
XX AAD22684:
XX
XX 26-FEB-2002 (first entry)
DE Streptomycetes fradiiae S-adenosylmethionine (SAM) operon DNA.
XX
XX K-Sadenosylmethionine; SAM operon; SAM synthetase; methyltransferase;
KW mtr; methylene tetrahydrofolate reductase; MTRH; activated methyl cycle;
KM tylosin production; ds.
XX
OS Streptomycetes fradiiae.
XX
FH Key Location/Qualifiers
EH CDS          986..2209
FT             /*tag= a
FT             /product= "SAM_synthetase_protein"
FT             2241..3341
FT             /*tag= b
FT             /product= "methyltransferase protein"
FT             /transl_except= (2241..2243, aa:mec)
FT             3348..4255
FT             /*tag= c
FT             /product= "Methylene tetrahydrofolate reductase protein"
FT             /transl_except= (3348..3340, aa:mec)
PN US6312920-B1.
XX
XX PD 06-NOV-2001.
XX PF 22-OCT-1997; 97US-0955957.
XX PR 13-NOV-1996; 96US-030898P.
XX
XX (ELIL ) LILLY & CO ELI.
```







(ITOL)/ITO L Y.  
PA  
PA (SHERM/) SHERMAN B K.

Lajgudi RV, Ito LY, Sherman BK;  
PI  
DR WPl: 2002-163647/21.

Noval purified corn tassell-derived polynucleotide useful for  
PT determining altered gene expression, to recover regulatory elements and  
PT to follow inheritance of desirable characteristics through hybrid  
PT breeding programs -

Claim 1; SEQ ID 1485; 201pp; English.

The present sequence describes a purified corn tassell-derived  
CC polynucleotide sequence (cdps) comprising a nucleic acid sequence  
CC selected from those given in ABL/0627 to ABL/76933. The cdps sequences  
CC encode corn tassell-derived polypeptides (CDPs). The cdps sequences (I)  
CC can be used for determining altered gene expression, to recover  
CC regulatory elements and to follow inheritance of desirable  
CC characteristics through hybrid breeding programs. (II are also useful  
CC in the evaluation, and alteration of desired characteristics associated  
CC with growth and development, disease resistance, environmental  
CC adaptability, quality and yield, and as molecular markers for studying  
CC inheritance of multigene traits in a plant breeding program. (III can be  
used to produce a tassell-specific profile of gene transcription), a  
transcript image, to clone regulatory elements for use in transformation  
vectors, to express a polypeptide, to identify, isolate or extend  
identical or related corn tassell nucleic acid sequences from DNA  
libraries, in nucleic acid hybridisation or amplification technologies,  
as query sequences to determine homology of known sequences, as probe  
for use in Southern or Northern hybridisation, and to identify the  
presence of and/or to determine the degree of similarity between two  
(or more) nucleic acid sequences.

Sequence 297 BP; 58 A; 85 C; 77 G; 50 T; 27 other;

Query Match      15.9%; Score 37.4; DB 24; Length 297;  
Best Local Similarity    78.6%; Pred. No. 0.0076;  
Matches     44; Conservative    0; Mismatches    12; Indels        0; Gaps            0;

OY    161 GGTTATCATCAAGCGGCCGTCATACGGCCTATTGGCGGTGAGCAGCGCCGCCTTACC    216  
| | | | | | | | | | | | | | | | | | | | |  
Db    242 GCCTACTCAAAGACGGCGGCGCTAAGSCNACTTGGAAGGACACCCCTGACTTACCC    297

RESULT 27

AAA81476  
ID AAA81476 standard; DNA; 56485 BP.

AAAB1476;  
AC  
XX  
DT  
DE  
XX  
XX  
X N. meningitidis partial DNA sequence gmm\_24 SEQ ID NO:24.

X X  
KW Neisseria meningitidis; Neisseria gonorrhoeae; genome; immunogenic;  
KM antigen; vaccine; diagnosis; infection; antibacterial; identification;  
RW Meningococcus B; Memb; ds.  
XX  
OS  
PN  
XX  
WO200022430-A2.

PD  
PD 20-APR-2000.

PF  
PE 08-OCT-1999;    99WO-US23573.

PR 09-OCT-1998;    98US-0103794.

PR 30-APR-1999;    99US-0132068.

XA  
XA (CHIR ) CHIRON CORP.

Pt Frazer CM, Hickey E, Peterson J, Tettein H, Venter JC; Scarlati V;  
Pi Massignani V, Galeotti C, Mora W, Ratil G, Scarselli M,  
P1 Rappunni R, Pizsa M.  
xx  
xx WPI: 2000-318079/27.

Dt Isolated nucleotide sequences of *Neisseria meningitidis* which can be  
Pt used in the diagnosis and treatment of N.meningitidis infection and  
PT other Neisserial infections, for example, N.gonorrhoeae -  
xx  
PS Claim 7: Page 507-524; 1760pp: English.

CC The present invention describes methods of obtaining immunogenic  
CC proteins from *Neisseria* genomic sequences. AAA81453 to AAA82414  
CC represent specifically claimed *Neisseria meningitidis* genomic DNA  
CC sequences; AAA81260 to AAA81303 and AAB25620 to AAB25663 represent  
CC *Neisseria* DNA sequences and their corresponding proteins; AAA81254 to  
CC AAA81259 and AAA81304 to AAA81321 represent PCR primers used in the  
CC isolation of *Neisseria meningitidis* DNA sequences; and AAA81322 to  
CC AAA81452 represent *Neisseria meningitidis* MenB polynucleotide ORF  
CC sequences, which are all used in the exemplification of the present  
CC invention. The nucleic acid sequences, protein sequences, and antibodies  
CC against them, can be used in the manufacture of a composition. The  
CC composition can be used as a medicament (or in the manufacture of a  
CC medicament) for treating, preventing or diagnosing infection due to  
CC Neisserial bacteria. For example, some of the identified proteins could  
CC be components of vaccines against Meningococcus B; against all serotypes;  
CC and/or against all pathogenic *Neisseriae*. Identification of sequences  
CC from the bacterium will also facilitate production of biological probes,  
CC particularly organism-specific probes. Attempts to make efficacious  
CC Meningococcus B vaccines have failed mainly due to antigen tolerance.  
CC Multivalent vaccines have also been tried but none have successfully  
CC overcome antigenic variability. The provision of further, complete  
CC sequences may provide an opportunity to identify secreted or surface  
CC exposed proteins that may be presumed targets for the immune system and  
CC which are not antigenically variable or at least more conserved than  
CC other more variable regions.  
XX  
SQ Sequence 56485 BP; 12504 A; 14247 C; 16158 G; 13573 T; 3 other;

Query Match 14.7%; Score 34.6; DB 21; Length 56485;  
Best Local Similarity 44.5%; Pred. No. 0.67;  
Matches 49; Conservative 0; Mismatches 61; Indels 0; Gaps 0;

Oy . 113 ACGTACGTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNCAAGTTCATCAGA 172  
Db 54015 ACCCTCGCCCCCAAAGCATGCACAATGCTCATCTCTTGCGCCCATTACAGTAAT 54074  
  
Oy 173 CGCGGGCAATAGGCCACTTTGGCCGTGACGAGCCGCAGCTCACCTCGCAG 222  
Db 54075 CGCGCGCTTAGCGACATTTGCGCCGCGAAGAACCTGACTTACCTGGAG 54124

RESULT 28  
AAE21612/C  
ID AAE21612 standard; DNA: 349980 BP.

XX  
XX AA#21612;  
XX  
DT 13-MAR-2001 (first entry)  
XX  
DE *Neisseria meningitidis* B nucleotide sequence SEQ ID NO:113.  
XX  
XX  
KW *Neisseria meningitidis*; *Neisseria gonorrhoeae*; immunogenic; vaccine;  
RV diagnosis; antigen; detection; infection; gene therapy; antibacterial;  
ds.  
XX  
OS *Neisseria meningitidis*.

PD 09-NOV-2000.

[illegible]

XX		
PN	WO200022430-A2.	
XX		
PD	20-APR-2000.	
XX		
PF	08-OCT-1999; 99WO-US23573.	
XX		
PR	09-OCT-1998; 98US-0103794.	
PR	30-APR-1999; 99US-0132068.	
PA	(CHIR) CHIRON CORP.	
PI	Frazier CM, Hickey E, Peterson J, Tetteijn H, Venter JC;	
PI	Masignani V, Galeotti C, Mora M, Ratti G, Scarselli M, Scarlato V;	
PI	Rappuoli R, Pizzo M;	
DR	WPI: 2000-318079/27.	
XX		
PT	Isolated nucleotide sequences of Neisseria meningitidis which can be	
PT	used in the diagnosis and treatment of N. meningitidis infection and	
PT	other Neisserial infections, for example, N.gonorrhoea -	
XX		
PS	Claim 7; Page 629-865; 1760pp; English.	
XX		
CC	The present invention describes methods of obtaining immunogenic	
CC	proteins from Neisseria genomic sequences. AAA81453 to AAA82414	
CC	represent specifically claimed Neisseria meningitidis genomic DNA	
CC	sequences: AAA81260 to AAA81303 and AAB25620 to AAB25663 represent	
CC	Neisseria DNA sequences and their corresponding proteins: AAA81254 to	
CC	AAA81259 and AAA81304 to AAA81321 represent PCR primers used in the	
CC	isolation of Neisseria meningitidis DNA sequences; and AAA81322 to	
CC	AAA81452 represent Neisseria meningitidis MenB polynucleotide ORF	
CC	sequences, which are all used in the exemplification of the present	
CC	invention. The nucleic acid sequences, protein sequences, and antibodies	
CC	against them, can be used in the manufacture of a composition. The	
CC	composition can be used as a medicament (or in the manufacture of a	
CC	medicament) for treating, preventing or diagnosing infection due to	
CC	Neisserial bacteria. For example, some of the identified proteins could	
CC	be components of vaccines against Meningococcus B; against all serotypes;	
CC	and/or against all pathogenic Neissariae. Identification of sequences	
CC	from the bacterium will also facilitate production of biological probes,	
CC	particularly organism-specific probes. Attempts to make efficacious	
CC	Meningococcus B vaccines have failed mainly due to antigen tolerance.	
CC	Multivalent vaccines have also been tried but none have successfully	
CC	overcome antigenic variability. The provision of further, complete	
CC	sequences may provide an opportunity to identify secreted or surface	
CC	exposed proteins that may be presumed targets for the immune system and	
CC	which are not antigenically variable or at least more conserved than	
CC	other more variable regions.	
SX		
S0	Sequence 837096 BP; 207534 A; 227065 C; 205215 G; 197280 T; 2 other:	
	Query Match 14.7%; Score 34.6; DB 21; Length 837096;	
	Best Local Similarity 44.5%; Pred.No. 2.1;	
	Matches 49; Conservative 0; Mismatches 61; Indels 0; Gaps 0;	
QY	113 ACGTACGTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNCAAGTCATCACAAGA 172	
D0	452410 ACCTTGCGGCCCAAAGGCATCGTCCAATGCTGTGATCTCTTGCGGCCGATTACAGTAAT 452351	
OY	173 CGCGCCGATACGGCCACTTTGGCCGTGAAGCACCGCCAGCTTCACTGCGAG 722	
D0	452350 CGCGCGCTTAGGACCATTTGCGCGCGCAAGAACCTGAGATTCACTTGGGAG 452301	
RESULT 30		
ID	AAI99683	
XX	AAI99683 standard; DNA: 4403765 BP.	
XX	AAI99683;	
XX		
DT	15-JAN-2002 (first entry)	
XX		











[illegible]

DB 85 CCGTGGCTGCGCCGCCGTTTTCACAGCCGCCGCTACTGTGTACGTAAACATGCGGGGTAC 144

OY 184 GGCCACTTTGGCCGTCAGCAGCGCATTTCACTTGACGAGTGTCGACACC 233  
||||| ||||| | ||||| ||| | |||||  
Db 145 CGCCAGATTGGCGCTCAATAACGACGATCTCCTCGAGGAGAGAACGAGAC 194

RESULT 39  
AAV64373  
ID AAV64373 standard; cDNA: 1519 BP.  
XX AAV64373;  
AC  
DT 15-FEB-1999 (first entry)  
XX  
DE GABA-gated chloride channel TBW-a3 cDNA.  
XX  
KM TBW-a3; GABA-gated chloride channel; tobacco budworm; insecticide;  
KM SS.  
XX  
OS Heliothis virescens.  
XX  
PH Key Location/Qualifiers  
FH CDS 1..1446  
FT /\*tag= a  
FT slg-peptide 1..66  
FT /\*tag= b  
FT slg-peptide 67..1443  
FT /\*tag= c  
XX  
XX WO9849185-A1.  
XX  
XX PD 05-NOV-1998.  
XX  
XX PF 27-APR-1998; 98WO-US08563.  
XX  
PR 02-JAN-1998; 98US-0002361.  
PR 28-APR-1997; 97US-0044976.  
PA (FMCC ) FMC CORP.  
PA  
PI Halling BP, Yuhas DA;  
PI  
XX WPI: 1999-009411/01.  
DR P-PSDB: AAM81635-36.  
DR  
XX  
XX PT New isolated lepidoptera GABA-gated chloride channels - comprise 3  
PT isoforms isolated from the tobacco budworm *Heliothis virescens*, used  
PT for characterizing bioactive agents, e.g. insecticides  
PS  
PS Claim Id: Fig 2; 55pp; English.

This cDNA sequence encompasses the open reading frame encoding  
GABA-gated chloride channel TBW-a3 (see AAM81635-36) of tobacco  
budworm (*Heliothis virescens*). TBW-a3, TBW-a2 (see AAM81633-34)  
and TBW-a1 (see AAM81637) proteins are 3 receptor isoforms that show  
sequence homology to each other and to other insect GABA-gated  
chloride channels. TBW-2a cDNA was obtained from *H. virescens* 4th  
instar larva RNA by PCR and RACE amplifications. The invention  
provides expression vectors in which a nucleic acid encoding a  
GABA-gated chloride channel is driven by an inducible promoter, and  
a claimed process for producing a GABA-gated chloride channel by  
transformed cells. The GABA-gated channels or cells expressing  
them can be used for characterizing a bioactive agent (claimed),  
e.g. for use as an insecticide. Probes and primers that identify  
or amplify GABA-gated chloride channel nucleic acids of the  
invention are also claimed.

Sequence 1519 BP; 421 A; 362 C; 347 G; 388 T; 1 other:

Query Match 12.7%; Score 29.8; DB 20; Length 1519;  
Best Local Similarity 40.5%; Pred. No. 6.3;  
Matches 64; Conservative 0; Mismatches 94; Indels 0; Gaps 0;

Qy	142	NNNNNNNNNNNNNNN	CAGGTCATCAACCGCCGCATAGGCCACTTGGCCGAC	201
		" " " " " " " "	" " " " " " " "	
Dd	2167	TACTTCAGCATTAAGA	TCAGCTTTGGGCAAGGCTGGGGACCGCAGCTACAAAAGACAGAT	2228
Qy	202	GAGCGGAGCTTCACCTG	CGAAGGTGG	226
		" " " " " " " "	" " " " " " " "	
Dd	2227	ATCATGGGCTGCCCC	TCTGCTGG	2251

RESULT 41  
AAT86246/C  
D AAT86246 standard; cDNA to mRNA, 1683 BP.

ID	Accession	Strand	Gene	Product	Location/Qualifiers
AT86246/c	AT86246	standard	CDNA to mRNA	1683 BP	
AT86246	AT86246				
07-JAN-1998			(first entry)		
CDNA encoding mugwort pollen co-factor-independent phosphoglycerate mutase isoform At17.					
Cofactor-Independent phosphoglycerate mutase; PGM-1; E.C. 5.4.21; Mugwort; pollen; allergy; plant allergen; panallergen; B cell; T cell; epithel; immunotherapy; detection; diagnosis; hay fever; conserved; ds.					
Artemisia vulgaris.					
Key	Location/Qualifiers				
CDS	1..1683				
	/tag= a				
MO9705258-A2.					
13-FEB-1997.					
02-AUG-1996;	96MO-AT00141.				
02-AUG-1995;	95AT-0001320.				
(BIOM-) BIOMAT PRODN & HANDELS GMBH.					
Breitenbach M, Ebner C, Engel E, Ferreira F, Jilek A; Kraft D, Richter K, Rheinberger H;					
MP1: 1997-145695/13.					
P-PSDB; AAM28505.					
New recombinant DNA encoding plant phosphoglycerate mutase or its antigenic epitope(s) - useful for diagnosis or treatment of allergies to pollen and plant-derived foods					
Claim 1: Fig 10b; 160pp; German.					
AT86246 encodes mugwort pollen co-factor-independent phosphoglycerate mutase (PGM-1) isoform At17. PGM-1 is a highly conserved plant allergen (panallergen) which can cause cross-reactivity in patients allergic to pollen and plant-derived foods. PGM-1 and it's B cell and T cell epitopes can be used for the in vitro detection of allergy against PGM-1, by measuring serum IgE or a cellular reaction. They can also be used in immunotherapy and will not cause an autoimmune response because PGM-1 is significantly different from the human enzyme, which is co-factor dependent.					
Sequence 1683 BP; 466 A; 304 C; 456 G; 457 T; 0 other;					

Query Match:	12.5%	Score	29.4	DB	18;
Best Local Similarity:	40.9%	Pred.	No. 9.1;		
Matches	54;	Conservative	0;	Mismatches	78; Indels 0; Gaps 0;
Oy	98	TGAGCAGGCACACACACTCTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNN	157		
bob	1421	TGTATCGCAGTGTCACACAATAATTTCCACTACTCTTAATTGCATCAAGATCATCTTA	1362		



SEQ Sequence 2604 BP; 779 A; 515 C; 579 G; 731 T; 0 other;

Query Match 12.4%; Score 29.2; DB 21; Length 2604;  
Best Local Similarity 75.8%; Pred. No. 13;  
Matches 50; Conservative 0; Mismatches 13; Indels 3; Gaps 1;

OY 38 CCATGCTGATAATGACGCTCTGATTCATTTGTTGTTATTAATGTGATTAAT 97  
DB 2330 CCATGCTGATAATGATGGCTCTGTTCCCTTT---TGTTTATTAATGTTGATTAAT 2386  
OY 98 TGAGCA 103  
DB 2387 TGAGCA 2392

RESULT 44  
ABLO4431/C  
ID ABLO4431 standard; cDNA; 1086 BP.

AC ABLO4431;  
XX  
XX 26-MAR-2002 (first entry)  
DT  
XX Drosophila melanogaster expressed polynucleotide SEQ ID NO 7775.  
DE  
XX Drosophila; developmental biology; cell signalling; insecticide;  
KW pharmaceutical; gene; ss.  
XX  
OS Drosophila melanogaster.  
XX  
PN WO200171042-A2.  
XX  
XX 27-SEP-2001.  
PD  
XX 23-MAR-2001; 2001WO-US09231.  
XX  
XX 23-MAR-2000; 2000US-191637P.  
PR 11-JUL-2000; 2000US-0614150.  
XX  
XX (PERK ) PE CORP NY.  
PA  
PI Venter JC, Adams M, Li PMD, Myers EW;  
XX  
XX WPI: 2001-656660/75.  
DR P-PSDB: ABB60328.  
XX  
XX New isolated nucleic acid detection reagent for detecting 1000 or more  
PT genes from Drosophila and for elucidating cell signalling and cell-cell  
PT interactions -  
XX  
XX Claim 1; SEQ ID NO 7775; 21bp + Sequence Listing; English.  
PS  
XX The invention relates to an isolated nucleic acid detection reagent  
CC capable of detecting 1000 or more genes from Drosophila. The invention is  
CC useful in developmental biology and in elucidating cell signalling and  
CC cell-cell interactions in higher eukaryotes for the development of  
CC insecticides, therapeutics and pharmaceutical drugs. The invention  
CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA  
CC sequences (ABR57737-ABR72072) and the encoded proteins  
CC (ABR57737-ABR72072).  
CC The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX  
XX Sequence 1086 BP; 295 A; 267 C; 273 G; 251 T; 0 other;

Query Match 12.3%; Score 29; DB 23; Length 1086;  
Best Local Similarity 40.0%; Pred. No. 10;  
Matches 68; Conservative 0; Mismatches 102; Indels 0; Gaps 0;

OY 36 GCCATGCTGATAAATGACGCTCTGATTCATTTGTTGTTATTAATGTGATTAAT 95  
DB 943 GCCATGCTGATAAATGACGCTCTGATTCATTTGTTGTTATTAATGTGATTAAT 884

OY 96 ATTGAGCAGACACACACAGCTAGCTTTNNNNNNNNNNNNNNNNNNNNNNNNNNNN 155  
DB 883 ATTGATTCACACACCTGCGCACCAAGCATCAGCGGGCTACCGAATCTCCAGTCGG 824  
OY 156 NNNCAGGTTGATCAAGACCGCCGATACGCGCACTTTGGCGTGACGAG 205  
DB 823 TCTCGTGCCATCCATCCATCCATGCGACATCATGATGATGCTCTTATTCGACG 774

RESULT 45  
AAH66738  
ID AAH66738 standard; DNA; 1221 BP.

AC AAH66738;  
XX  
XX 26-SEP-2001 (first entry)  
DT  
XX C glutamicum coding sequence fragment SEQ ID NO: 1773.  
DE  
XX Corynebacterium; amino acid synthesis; vitamin; saccharide;  
KW organic acid synthesis; ds.  
XX  
OS Corynebacterium glutamicum.  
XX  
XX EP1108790-A2.  
PN  
XX 20-JUN-2001.  
PD  
XX 18-DEC-2000; 2000EP-0127688.  
PF  
XX 16-DEC-1999; 99JP-0377484.  
PR 07-APR-2000; 2000JP-0159162.  
PR 03-AUG-2000; 2000JP-0280988.  
XX  
XX (KJOM ) KYOMA HAKKO KOGYO KK.  
PA  
XX Nakagawa S, Mizoguchi H, Ando S, Hayashi M, Ochiai K, Yokoi H;  
PI Tateishi N, Senoh A, Ikeda M, Ozaki A;  
XX  
XX WPI: 2001-376931/40.  
DR P-PSDB: AAG91519.  
XX  
XX Novel polynucleotides derived from Corynebacterium bacteria, for identifying  
PT mutation point of a gene, measuring expression of a gene, analysing  
PT expression profile or pattern of a gene and identifying homologous gene  
PT -  
XX  
XX Claim 8; SEQ ID NO: 1773; 246bp + Sequence Listing; English.  
PS  
XX The present invention provides a number of nucleotide and protein  
CC sequences from the Corynebacterium bacterium Corynebacterium glutamicum. These  
CC are useful for identifying the mutation point of a gene derived from a  
CC mutant of corynebacterium bacterium, measuring expression amount and  
CC analysing the expression profile or expression pattern of a gene derived  
CC from Corynebacterium bacterium, and identifying a homologue of a gene derived  
CC from corynebacterium bacterium. Corynebacterium bacteria are useful for producing  
CC amino acids, nucleic acids, vitamins, saccharides and organic acids,  
CC particularly L-lysine. The present sequence is a nucleic acid described  
CC in the exemplification of the invention.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from the  
CC European Patent Office.  
XX  
XX Sequence 1221 BP; 272 A; 350 C; 336 G; 263 T; 0 other;

Query Match 12.3%; Score 29; DB 22; Length 1221;  
Best Local Similarity 41.0%; Pred. No. 11;  
Matches 50; Conservative 0; Mismatches 72; Indels 0; Gaps 0;

OY 113 ACGTCAGTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 172  
DB 1067 ACGTCAGTCAGCAGCAATTAATCCGTGAGTTGATTCGCTTCGTCGATCTACGCTGACA 1126

Wed Apr 23 15:43:04 2003

us-09-198-779b-1.rng

Page 33

OY 173 CCGGCGATTACGGCCACTTTTGGCCGTGAGACGCCGCATCTTCACCTGCAGGTGTCACG 232  
| | | | | | | | | | | | | | | | | |  
Db 1127 CTGCTGCTACGGCCACTTTGGTGCGCATGTAITTTGACCTTCTTTGGAGGCTATCGACC 1186  
  
OY 233 CC 234  
|  
Db 1187 GC 1188

Search completed: April 23, 2003, 15:01:42  
Job time : 3205.99 secs

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GenCore version 5.1.4-p5.4578  
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## OM nucleic - protein search, using frame\_plus\_n2p model

Run on: April 23, 2003, 12:06:28 : Search time 19.5833 Seconds  
(without alignments)  
2307.226 Million cell updates/sec

Title: US-09-198-779b-1

Perfect score: 349  
Sequence: 1 gtttcgcgtctagctcgtt.....ctgcgaggtggtcgaagccccc 235

Scoring table:

BLOSUM62  
Xgapop 10.0, Xgapext 0.5  
Ygapop 10.0, Ygapext 0.5  
Fgapop 6.0, Fgapext 7.0  
Delop 6.0, Delext 7.0

Searched: 283224 seqs, 96134422 residues

Total number of hits satisfying chosen parameters: 566448

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 50 summaries

Command line parameters:  
-MODEL=frame+n2p.model -DEV=xip  
-O=/cgn2.1/USPTO.spool/US09198779.r/unat.23042003.120617.2416/app.query.fasta.1.917  
-DB=PIR.73 -OFT=fastan -SUFFIX=rrr -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0  
-UNITS=bites -START=1 -END=1 -MATRIX=blonum62 -TRANS=human40.cdi -LIST=50  
-DOCALLIGN=200 -THR\_SCORE=pct -THR\_MAX=100 -THR\_MIN=0 -ALIGN=45 -MODE=LOCAL  
-OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000  
-USER=US09198779.qcgn.1.1\_92@unat.23042003.120617.2416 -NCPU=6 -ICPU=3  
-NO\_XLPHY -NO\_MMAPP -LARGEQUERY -NEG\_SCORES=0 -WAIT -LONGLOG -DEV.TIMEOUT=120  
-WARN.TIMEOUT=30 -THREDS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6 -Fgapext=7  
-YGAPOP=10 -YGAPEXT=0.5 -DELop=6 -DELEXT=7

Database: PIR.73:\*

1: pir1:\*\n2: pir2:\*\n3: pir3:\*\n4: pir4:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	127	36.4	394	2	T06180 methionine adenosyl
2	120	34.4	68	2	PQ0817 methionine adenosyl
3	120	34.4	393	2	JN0131 methionine adenosyl
4	120	34.4	393	2	S38875 methionine adenosyl
5	120	34.4	393	2	S46538 methionine adenosyl
6	120	34.4	393	2	S46538 methionine adenosyl
7	116	33.2	360	2	T06582 methionine adenosyl
8	116	33.2	366	2	S66351 methionine adenosyl
9	115	33.0	390	2	G84785 probable s-adenosyl
10	115	33.0	393	2	JQ0410 methionine adenosyl
11	108.5	31.1	397	2	S66352 methionine adenosyl
12	101	28.9	179	2	T07899 methionine adenosyl
13	101	28.9	390	2	S49491 methionine adenosyl
14	101	28.9	390	2	S46540 methionine adenosyl

15	98	28.1	396	2	T10710 methionine adenosyl
16	74	21.2	395	2	G72228 S-adenosylmethionin
17	72	20.6	389	2	E81986 probable methionin
18	72	20.6	389	2	D81042 S-adenosylmethionin
19	68	19.5	388	2	S51671 methionine adenosyl
20	65	18.6	400	2	D69657 methionine adenosyl
21	64.5	18.5	407	2	B87255 S-adenosylmethionin
22	64	18.3	376	2	A82895 S-adenosylmethionin
23	63.5	18.4	1979	2	JW0059 mtrpD protein - mo
24	63	18.1	395	1	S27257 methionine adenosyl
25	63	18.1	395	2	A37118 methionine adenosyl
26	63	18.1	403	2	H86976 probable S-adenosyl
27	63	18.1	589	2	G87485 hypothetical prote
28	61	17.5	376	2	F90589 S-adenosylmethionin
29	61	17.5	385	2	D71964 S-adenosylmethionin
30	61	17.5	385	2	E64544 methionine adenosyl
31	61	17.5	398	2	F89664 S-adenosylmethionin
32	61	17.5	399	2	D84062 S-adenosylmethionin
33	61	17.5	399	2	F86862 methionine adenosyl
34	61	17.5	399	2	AD1654 S-methionine adeno
35	61	17.5	403	2	F70899 probable metk prot
36	61	17.5	403	2	B97403 methionine adenosyl
37	61	17.5	420	2	AB2621 S-adenosylmethionin
38	61	17.5	420	2	AB2621 heme transport pro
39	61	17.5	877	2	AC2211 methionine adenosyl
40	60.5	17.3	409	2	S74736 S-adenosylmethionin
41	60	17.2	385	2	E82319 S-adenosylmethionin
42	60	17.3	674	2	B84381 acylaminoacyl-pept
43	59	16.9	396	2	A71281 probable S-adenosyl
44	59	16.9	396	2	A47151 methionine adenosyl
45	59	16.9	397	2	S06114 hypothetical prote
46	58.5	16.9	155	2	H89787 hypothetical prote
47	58.5	16.8	238	2	H70866 methionine adenosyl
48	58	16.6	378	2	E84977 ribosomal protein
49	58	16.6	472	2	G75298 microtubule-associ
50	58	16.6	1861	2	T13845

## ALIGNMENTS

## RESULT 1

T06180 methionine adenosyltransferase (EC 2.5.1.6) - barley

N:Alternate names: S-adenosylmethionine synthetase

C:Species: Hordeum vulgare (barley)

C:Date: 30-Apr-1999 #sequence\_revision 30-Apr-1999 #text\_change 20-Jun-2000

C:Accession: T06180

R:Mori, S. submitted to the EMBL Data Library, August 1995

A:Reference number: Z15512

A:Accession: T06180

A:Status: translated from GB/EMBL/DBJ

A:Molecule type: mRNA

A:Residues: 1-394 <MOR>

A:Cross-references: EMBL:D63835; PIDN:BAA0995.1

C:Superfamily: methionine adenosyltransferase

C:Keywords: S-adenosylmethionine; transferase

## Alignment Scores:

Pred. No.: 2.4e-10 Length: 394  
Score: 127.00 Matches: 24  
Percent Similarity: 96.00% Conservative: 0  
Best Local Similarity: 96.00% Mismatches: 1  
Query Match: 36.39% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779b-1 (1-235) x T06180 (1-394)

OY AGGTTTCATCAGACCGGCATACGGCCATTGGCCGTGACGAGCCGACCTTCACCTGC 219  
|||||  
Db 362 ArghpellelystmrAlaAlaTyrClyHisPhcglyArGAspAlaAspPheThrTrp 381  
OY 220 GAGGTGTCAGACCCC 234

Db 382 GluValVallylPro 386

# RESULT 2

C:Species: Brassica napus (rape)  
C:Date: 03-May-1994 #sequence\_revision 07-Oct-1994 #text\_change 05-May-2000

C:Accession: PQ0817  
R:Park, Y.S.; Kwak, J.M.; Kwon, O.Y.; Kim, Y.S.; Lee, D.S.; Cho, M.J.; Lee, H.H.; Nam, H.

Plant Physiol. 103, 359-370, 1993  
A:Title: Generation of expressed sequence tags of random root cDNA clones of Brassica na

A:Reference number: PQ0816; MWID:94302145; PMID:8029352

A:Accession: PQ0817  
A:Molecule type: mRNA

A:Residues: 1-68 <PAR>  
A:Experimental source: root, cv. Naehan

C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase

## Alignment Scores:

Pred. No.:	2,54e-09	Length:	68
Score:	120.00	Matches:	22
Percent Similarity:	92.00%	Conservative:	1
Best Local Similarity:	88.00%	Mismatches:	2
Query Match:	34.38%	Indels:	0
DB:	2	Gaps:	0

US-09-198-779B-1 (1-235) x PQ0817 (1-68)

QY 160 AGGTTTCATCAAGACCGCGCATAGCGCCACTTTGGCCGTGACGACGCCGACTTCACCTGC 219

Db 36 ArgPheLeuysThrAlaAlaTyGlyHisPheGlyArgAspAspProAspPheThrTrp 55

QY 220 GAGGTGTCACAGCCC 234

Db 56 GluValVallylPro 60

# RESULT 3

methionine adenosyltransferase (EC 2.5.1.6) - Arabidopsis thaliana  
N:Alternate names: S-adenosylmethionine synthetase

C:Species: Arabidopsis thaliana (mouse-ear cress)  
C:Date: 30-Sep-1991 #sequence\_revision 30-Sep-1991 #text\_change 05-May-2000

C:Accession: JN0131  
R:Peleman, J.; Boerjan, W.; Engler, G.; Seurinck, J.; Botterman, J.; Alliotte, T.; Van M

Plant Cell 1, 81-93, 1989  
A:Title: Strong cellular preference in the expression of a housekeeping gene of Arabidop

A:Reference number: JN0131; MWID:92386056; PMID:2535470

A:Accession: JN0131  
A:Molecule type: DNA

A:Residues: 1-393 <PEL>  
A:Cross-references: GB:M55077; NID:q166871; PIDN:AAA32866.1; PID:q166872

A:Experimental source: var. K85  
A:Note: the sequence derived from var. Columbia differs from that shown in having 117-Glu

C:Comment: S-Adenosylmethionine synthetase catalyzes the biosynthesis of adenosylmethio

C:Gene: sam-1  
C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase

## Alignment Scores:

Pred. No.:	2,76e-09	Length:	393
Score:	120.00	Matches:	22
Percent Similarity:	92.00%	Conservative:	1
Best Local Similarity:	88.00%	Mismatches:	2
Query Match:	34.38%	Indels:	0
DB:	2	Gaps:	0

US-09-198-779B-1 (1-235) x JN0131 (1-393)

QY 160 AGGTTTCATCAAGACCGCGCATAGCGCCACTTTGGCCGTGACGACGCCGCGACTTCACCTGC 219

Db 361 ArgPheLeuysThrAlaAlaTyGlyHisPheGlyArgAspAspProAspPheThrTrp 380

QY 220 GAGGTGTCACAGCCC 234

Db 381 GluValVallylPro 385

# RESULT 4

methionine adenosyltransferase (EC 2.5.1.6) - tomato  
N:Alternate names: S-adenosyl-L-methionine synthetase

C:Species: Lycopersicon esculentum (tomato)  
C:Date: 22-Jan-1994 #sequence\_revision 10-Nov-1995 #text\_change 05-May-2000

C:Accession: S46539; S38875  
R:Espartero, J.; Pintor-Toro, J.A.; Pardo, J.M.

Plant Mol. Biol. 25, 217-227, 1994  
A:Title: Differential accumulation of S-adenosylmethionine synthetase transcripts in

A:Reference number: S46538; MWID:94289646; PMID:8018871

A:Accession: S46539  
A:Status: nucleic acid sequence not shown

A:Residues: 1-393 <ES2>  
A:Cross-references: EMBL:Z24742; NID:q429105; PIDN:CAA80866.1; PID:q429106

C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase

## Alignment Scores:

Pred. No.:	2,76e-09	Length:	393
Score:	120.00	Matches:	22
Percent Similarity:	92.00%	Conservative:	1
Best Local Similarity:	88.00%	Mismatches:	2
Query Match:	34.38%	Indels:	0
DB:	2	Gaps:	0

US-09-198-779B-1 (1-235) x S38875 (1-393)

QY 160 AGGTTTCATCAAGACCGCGCATAGCGCCACTTTGGCCGTGACGACGCCGACTTCACCTGC 219

Db 361 ArgPheLeuysThrAlaAlaTyGlyHisPheGlyArgAspAspProAspPheThrTrp 380

QY 220 GAGGTGTCACAGCCC 234

Db 381 GluValVallylPro 385

# RESULT 5

methionine adenosyltransferase (EC 2.5.1.6) - tomato  
N:Alternate names: S-adenosyl-L-methionine synthetase

C:Species: Lycopersicon esculentum (tomato)  
C:Date: 26-Dec-1994 #sequence\_revision 10-Nov-1995 #text\_change 05-May-2000

C:Accession: S46538; S38874  
R:Espartero, J.; Pintor-Toro, J.A.; Pardo, J.M.

Plant Mol. Biol. 25, 217-227, 1994  
A:Title: Differential accumulation of S-adenosylmethionine synthetase transcripts in

A:Reference number: S46538; MWID:94289646; PMID:8018871

A:Accession: S46538  
A:Status: nucleic acid sequence not shown

A:Residues: 1-393 <ESP>  
A:Cross-references: EMBL:Z24741; NID:q429103; PIDN:CAA80865.1; PID:q429104

C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase

## Alignment Scores:

Pred. No.:	2,76e-09	Length:	393
Score:	120.00	Matches:	22
Percent Similarity:	92.00%	Conservative:	1
Best Local Similarity:	88.00%	Mismatches:	2
Query Match:	34.38%	Indels:	0
DB:	2	Gaps:	0

US-09-198-779B-1 (1-235) x S46538 (1-393)





R; Gomez, L.; Carrasco, P.

## RESULT 13

R; Gomez, L.; Carrasco, P.

methionine adenosyltransferase (EC 2.5.1.6) - garden petunia  
C:Species: Petunia x hybrida (garden petunia)  
C:Date: 01-Feb-1995 #sequence\_revision 10-Nov-1995 #text\_change 05-May-2000  
C:Accession: S49491  
R:Izhaki, A.; Shoseyov, O.; Weiss, D.  
submitted to the EMBL Data Library, October 1994  
A:Description: Petunia cDNA encoding S-Adenosylmethionine synthetase.  
A:Reference number: S49491  
A:Accession: S49491  
A:Molecule type: mRNA  
A:Residues: 1-390 <I2>  
A:Cross-references: EMBL:X82214; NID:q559505; PIDN:CA57696.1; PID:q559506  
C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
Pred. No.: 2.1e-06 Length: 390  
Score: 101.00 Matches: 19  
Percent Similarity: 83.33% Conservative: 1  
Best Local Similarity: 79.17% Mismatches: 4  
Query Match: 28.94% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1 (1-235) x S49491 (1-390)

OY 160 AGGTTTCATCAGACCGCGCATACGCGCCTTTGGCCGTGACGACCGCGACTTCACCTGC 219  
||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||  
DB 361 ArgTYrGlnLYsThrAlaAlaIaTyrgLYsHPheGLyArgSPaSProAspPhehrrTP 380  
OY 220 GAGGTGCTCAG 231  
||| |||||  
DB 381 GluThrValLYs 384

RESULT 14  
S46540  
methionine adenosyltransferase (EC 2.5.1.6) - tomato  
N:Alternate names: S-adenosyl-L-methionine synthetase  
C:Species: Lycopersicon esculentum (tomato)  
C:Date: 26-Dec-1994 #sequence\_revision 10-Nov-1995 #text\_change 05-May-2000  
C:Accession: S46540; S38876  
R:Espartero, J.; Plator-Toro, J.A.; Pardo, J.M.  
Plant Mol. Biol. 25, 217-227, 1994  
A:Title: Differential accumulation of S-adenosylmethionine synthetase transcripts in res  
A:Reference number: S46538; MUID:94289646; PMID:8018871  
A:Accession: S46540  
A:Status: nucleic acid sequence not shown  
A:Molecule type: mRNA  
A:Residues: 1-390 <ESP>  
A:Cross-references: EMBL:Z24743; NID:q429107; PIDN:CA80867.1; PID:q429108  
C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
Pred. No.: 2.1e-06 Length: 390  
Score: 101.00 Matches: 19  
Percent Similarity: 83.33% Conservative: 1  
Best Local Similarity: 79.17% Mismatches: 4  
Query Match: 28.94% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1 (1-235) x S46540 (1-390)

OY 160 AGGTTTCATCAGACCGCGCATACGCGCCTTTGGCCGTGACGACCGCGACTTCACCTGC 219  
||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||  
DB 361 ArgTYrGlnLYsThrAlaAlaIaTyrgLYsHPheGLyArgSPaSProAspPhehrrTP 380  
OY 220 GAGGTGCTCAG 231  
||| |||||  
DB 381 GluThrValLYs 384

RESULT 15  
T10710  
methionine adenosyltransferase (EC 2.5.1.6) - clove pink

N:Alternate names: S-adenosylmethionine synthetase  
C:Species: Dianthus caryophyllus (clove pink)  
C:Date: 16-Jul-1999 #sequence\_revision 16-Jul-1999 #text\_change 05-May-2000  
C:Accession: T10710  
R:Larsen, P.B.; Woodson, W.R.  
submitted to the EMBL Data Library, April 1991  
A:Description: Cloning and nucleotide sequence of a S-adenosylmethionine synthetase c  
A:Reference number: 217091  
A:Accession: T10710  
A:Status: translated from GB/EMBL/DBJ  
A:Molecule type: mRNA  
A:Residues: 1-396 <LAR>  
A:Cross-references: EMBL:M61882; NID:q167961; PID:q304637  
C:Genetics: SAM2  
A:Gene: SAM2  
C:Function: catalyzes the formation of S-adenosyl methionine with phosphate and py  
C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
Pred. No.: 5.98e-06 Length: 396  
Score: 98.00 Matches: 17  
Percent Similarity: 83.33% Conservative: 3  
Best Local Similarity: 70.83% Mismatches: 4  
Query Match: 28.08% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1 (1-235) x T10710 (1-396)

OY 160 AGGTTTCATCAGACCGCGCATACGCGCCTTTGGCCGTGACGACCGCGACTTCACCTGC 219  
||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||  
DB 364 ArgTYrLeuLYsThrAlaAlaIaTyrgLYsHPheGLyArgGLuAsPProAspPhehrrTP 383  
OY 220 GAGGTGCTCAG 231  
||| |||||  
DB 384 GluAlaAlaLYs 387

RESULT 16  
G72228  
S-adenosylmethionine synthetase - Thermotoga maritima (strain MSB8)  
C:Species: Thermotoga maritima  
C:Date: 11-Jun-1999 #sequence\_revision 11-Jun-1999 #text\_change 21-Jul-2000  
C:Accession: G72228  
R:Nelson, K.E.; Clayton, R.A.; Gill, S.R.; Gwin, M.L.; Dodson, R.J.; Haft, D.H.; Hic  
Garrett, M.M.; Stewart, A.M.; Cotton, M.D.; Pratt, M.S.; Phillips, C.A.; Richardson,  
C.M.  
Nature 399, 323-329, 1999  
A:Title: Evidence for lateral gene transfer between Archaea and Bacteria from genome  
A:Reference number: A72200; MUID:99287316; PMID:10360571  
A:Accession: G72228  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-395 <ARN>  
A:Cross-references: GB:AE001807; GB:AE000512; NID:q4982216; PIDN:AA036725.1; PID:q498  
A:Experimental source: strain MSB8  
C:Genetics: TML658  
A:Gene: TML658  
C:Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 0.026 Length: 395  
Score: 74.00 Matches: 13  
Percent Similarity: 88.89% Conservative: 3  
Best Local Similarity: 72.22% Mismatches: 2  
Query Match: 21.20% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1 (1-235) x G72228 (1-395)

OY 169 AAGACCGCGCGCATACGCGCCTTTGGCCGTGACGACCGCGACTTCACCTGCAG 222  
||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||  
DB 364 LysThrAlaAlaIaTyrgLYsHPheGLyArgAsnGLuAsnGLuPhehrrTP 381







```

Best Local Similarity: 31.15%      Mismatches: 27
Query Match: 18.05%      Indels: 12
DB: 2      Gaps: 2

US-09-198-779B-1 (1-235) x G87485 (1-589)

OY 53 GACGGTCCTGATTCATGTTGTTGTTATTAATGATGACGACACAC 112
    |||||
DB 240 AspGlyProAspPro-----GlnGlyProMet 248

OY 113 ACCTACCTTTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 172
    |||||
DB 249 ThrlslnAlaAlaAlaAlaAlaAlaAlaAlaAlaAlaAlaAla 267

OY 173 CCGCCGATAGCGCCACTTTGGCCGTGACGACGCCACTTACCTGCGAGG 232
    |||||
DB 268 ProProAlaGlnSerAlaArgLeuAlaAlaAlaAlaAlaAlaAla 287

OY 233 CCC 235
    |||
DB 288 Pro 288

RESULT 28
hypothetical protein MYPU_7020 [Imported] - Mycoplasma pulmonis (strain UAB CTIP)
C:Species: Mycoplasma pulmonis
C:Date: 24-May-2001 #sequence_revision 24-May-2001 #text_change 03-Aug-2001
C:Accession: F90599
R:Chamaud, I.; Hellig, R.; Ferris, S.; Barbe, V.; Sanson, D.; Gallsson, F.; Moszer, I.;
Nucleic Acids Res. 29, 2145-2153, 2001
A:Title: The complete genome sequence of the murine respiratory pathogen Mycoplasma pulm
A:Reference number: A99512; M0ID:21267165; PMID:11353084
A:Accession: F90599
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-376 <KUR>
A:Cross-references: GB:AL445566; PID:g14090117; PIDN:CAJ13875.1; GSPDB:GN00153
A:Experimental source: strain UAB CTIP
C:Genetics:
A:Gene: MYPU_7020
A:Genetic code: SGC3
C:Superfamily: methionine adenosyltransferase

Alignment Scores:
Pred. No.: 2.43      Length: 376
Score: 61.00      Matches: 10
Percent Similarity: 66.67%      Conservative: 4
Best Local Similarity: 47.62%      Mismatches: 7
Query Match: 17.48%      Indels: 0
DB: 2      Gaps: 0

US-09-198-779B-1 (1-235) x F90599 (1-376)

OY 160 AGGTTCAATCAAGACCGCCGATACGCCACTTTGGCCGTGACGACGCCGACTTACCTGC 219
    |||||
DB 343 LysTyrThrProThrSerPhePheGlyHisPheGlyArgAspSerLeuAspLeuProTrp 362

OY 220 GAG 222
    |||
DB 363 Glu 363

RESULT 29
D71964
s-adenosylmethionine synthetase - Helicobacter pylori (strain J99)
C:Species: Helicobacter pylori
A:Variety: strain J99
C:Date: 12-Feb-1999 #sequence_revision 12-Feb-1999 #text_change 18-Jun-1999
C:Accession: D71964
R:Alm, R.A.; Ling, L.S.L.; Moir, D.T.; King, B.L.; Brown, E.D.; Doig, P.C.; Smith, D.R.;
Ives, C.; Gibson, R.; Metberg, D.; Mills, S.D.; Jiang, Q.; Taylor, D.E.; Vovis, G.F.;
Nature 397, 176-180, 1999
A:Title: Genomic sequence comparison of two unrelated isolates of the human gastric pat
A:Reference number: A71800; M0ID:99120557; PMID:9923682

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```

A:Accession: D71964
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-385 <ARN>
A:Cross-references: GB:AE001456; GB:AE001439; MID:g4154689; PIDN:AAD05755.1; PID:g415
A:Experimental source: strain J99
C:Genetics:
A:Gene: metK
C:Superfamily: methionine adenosyltransferase

Alignment Scores:
Pred. No.: 2.43      Length: 385
Score: 61.00      Matches: 11
Percent Similarity: 82.35%      Conservative: 3
Best Local Similarity: 64.71%      Mismatches: 3
Query Match: 17.48%      Indels: 0
DB: 2      Gaps: 0

US-09-198-779B-1 (1-235) x D71964 (1-385)

OY 172 ACCGCCGATAGCGCCACTTTGGCCGTGACGACGCCGACTTACCTGCAG 222
    |||||
DB 355 ThrSerAlaTyrGlyHisPheGlyArgGluLeuGluGluPheTrpGlu 371

RESULT 30
E64544
methionine adenosyltransferase (EC 2.5.1.6) 2 - Helicobacter pylori (strain 26695)
C:Species: Helicobacter pylori
C:Date: 09-Aug-1997 #sequence_revision 09-Aug-1997 #text_change 05-May-2000
C:Accession: E64544
R:Tomb, J.F.; White, O.; Kerlavage, A.R.; Clayton, R.A.; Sutton, G.G.; Fleischmann, R
Peterson, S.; Loftus, B.; Richardson, D.; Dodson, R.; Khalak, H.G.; Glodok, A.; McKe
son, J.D.; Kelley, J.M.; Cotton, M.D.; Weidman, J.M.; Fujii, C.; Bowman, C.; Watthey,
Nature 388, 539-547, 1997
A:Authors: Wallin, E.; Hayes, W.S.; Borodovsky, M.; Karp, P.D.; Smith, H.O.; Fraser,
A:Title: The complete genome sequence of the gastric pathogen Helicobacter pylori.
A:Reference number: A64520; M0ID:97394467; PMID:9252185
A:Accession: E64544
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-385 <TOM>
A:Cross-references: GB:AE000540; GB:AE000511; MID:g2313287; PIDN:AAD07267.1; PID:g231
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:
Pred. No.: 2.43      Length: 385
Score: 61.00      Matches: 11
Percent Similarity: 82.35%      Conservative: 3
Best Local Similarity: 64.71%      Mismatches: 3
Query Match: 17.48%      Indels: 0
DB: 2      Gaps: 0

US-09-198-779B-1 (1-235) x E64544 (1-385)

OY 172 ACCGCCGATAGCGCCACTTTGGCCGTGACGACGCCGACTTACCTGCAG 222
    |||||
DB 355 ThrSerAlaTyrGlyHisPheGlyArgGluLeuGluGluPheTrpGlu 371

RESULT 31
F89964
s-adenosylmethionine synthetase [Imported] - Staphylococcus aureus (strain N315)
C:Species: Staphylococcus aureus
C:Date: 10-May-2001 #sequence_revision 10-May-2001 #text_change 22-Oct-2001
C:Accession: F89964
R:Kuroda, M.; Ohca, T.; Uchiyama, I.; Baba, T.; Yuzawa, H.; Kobayashi, I.; Cui, L.; O
ma, A.; Mizutani-Ui, Y.; Kobayashi, N.; Sawano, T.; Inoue, R.; Kaito, C.; Sekimizu, K
C.; Shiba, T.; Hattori, M.; Ogasawara, N.; Hayashi, H.; Hiratsuyu, K.
Lancet 357, 1225-1240, 2001
A:Title: Whole genome sequencing of methicillin-resistant Staphylococcus aureus.
A:Reference number: A89758; M0ID:21311952; PMID:11418146
A:Accession: F89964
A:Status: preliminary

```

A: Molecule type: DNA  
A: Residues: 1-398 <RUR>  
A: Cross-references: GB:BA000018; PID: g13701583; PIDN: BAB42876.1; GSPDB: GN00149  
A: Experimental source: strain N315  
C: Genetics:  
A: Gene: metK  
C: Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 2.43 Length: 398  
Score: 61.00 Matches: 11  
Percent Similarity: 72.22% Conservative: 2  
Best Local Similarity: 66.67% Mismatches: 5  
Query Match: 17.48% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1 (1-235) x F89964 (1-398)

OY 169 AAGACCGCCGATACGCGCATTGGCCGTGACGACGCGCGACTTCACCTGCGAG 222  
DB 367 GlnThrAlaAlaIatYrGlyHisPheGlyArgThrAspValGluLeuProTrpGlu 384

RESULT 32

DB4062

S: adenosylmethionine synthetase metK [imported] - Bacillus halodurans (strain C-125)  
C: Species: Bacillus halodurans  
C: Date: 01-Dec-2000 #sequence\_revision 01-Dec-2000 #text\_change 15-Jun-2001  
C: Accession: DB4062  
R: Takami, H.; Nakasone, K.; Takaki, Y.; Maeno, G.; Sasaki, R.; Masui, N.; Fujii, F.; Hirai  
Nucleic Acids Res. 28, 4317-4331, 2000  
A: Title: Complete genome sequence of the alkaliphilic bacterium Bacillus halodurans and  
A: Reference number: AB3650; MUID: 20512582; PMID: 11058132  
A: Accession: DB4062  
A: Status: preliminary  
A: Molecule type: DNA  
A: Residues: 1-399 <STO>  
A: Cross-references: GB:AP001518; GB:BA000004; NID: g10175792; PIDN: BAB07019.1; GSPDB: GN00  
A: Experimental source: strain C-125  
C: Genetics:  
A: Gene: metK  
C: Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 2.43 Length: 399  
Score: 61.00 Matches: 11  
Percent Similarity: 72.22% Conservative: 2  
Best Local Similarity: 61.11% Mismatches: 5  
Query Match: 17.48% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1 (1-235) x DB4062 (1-399)

OY 169 AAGACCGCCGATACGCGCATTGGCCGTGACGACGCGCGACTTCACCTGCGAG 222  
DB 370 GlnThrAlaAlaIatYrGlyHisPheGlyArgThrAspValGluLeuProTrpGlu 387

RESULT 33

F86862

methionine adenosyltransferase (EC 2.5.1.6) [imported] - Lactococcus lactis subsp. lacti  
N: Alternate names: S-adenosylmethionine synthetase  
C: Species: Lactococcus lactis subsp. lactis  
C: Date: 23-Mar-2001 #sequence\_revision 23-Mar-2001 #text\_change 03-Aug-2001  
C: Accession: F86862  
R: Bolotin, A.; Winkler, P.; Mauger, S.; Jallou, O.; Malarme, K.; Welssenbach, J.; Ehrlich  
Genome Res. 11, 731-753, 2001  
A: Title: The complete genome sequence of the lactic acid bacterium Lactococcus lactis ss  
A: Reference number: AB6625; MUID: 21235186; PMID: 11337471  
A: Accession: F86862  
A: Status: preliminary  
A: Molecule type: DNA  
A: Residues: 1-399 <STO>  
A: Cross-references: GB:AE005176; PID: g12724937; PIDN: AAK06000.1; GSPDB: GN00146  
A: Experimental source: strain IL403

C: Genetics:  
A: Gene: metK  
C: Superfamily: methionine adenosyltransferase  
C: Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
Pred. No.: 2.43 Length: 399  
Score: 61.00 Matches: 11  
Percent Similarity: 72.22% Conservative: 2  
Best Local Similarity: 61.11% Mismatches: 5  
Query Match: 17.48% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1 (1-235) x F86862 (1-399)

OY 169 AAGACCGCCGATACGCGCATTGGCCGTGACGACGCGCGACTTCACCTGCGAG 222  
DB 368 GlnThrAlaAlaIatYrGlyHisPheGlyArgSerAspLeuAspLeuProTrpGlu 385

RESULT 34

AD1654

S: methionine adenosyltransferase homolog metK [imported] - Listeria innocua (strain C  
C: Species: Listeria innocua  
C: Date: 27-Nov-2001 #sequence\_revision 27-Nov-2001 #text\_change 14-Dec-2001  
C: Accession: AD1654  
R: Glaaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloec  
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihl,  
D.; Jones, L.M.; Karst, U.  
Science 294, 849-852, 2001  
A: Authors: Kreft, J.; Kuhn, M.; Kunst, F.; Kurapkai, G.; Madueno, E.; Maitournam, A.;  
Ok, C.; Schlueter, T.; Sinoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehla  
A: Title: Comparative genomics of Listeria species.  
A: Reference number: AB1077; MUID: 21537279; PMID: 11679669  
A: Accession: AD1654  
A: Status: preliminary  
A: Molecule type: DNA  
A: Residues: 1-399 <GLA>  
A: Cross-references: GB:AL592022; PIDN: CAC97004.1; PID: g16414260; GSPDB: GN00178  
A: Experimental source: strain C1P11262  
C: Genetics:  
A: Gene: metK  
C: Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 2.43 Length: 399  
Score: 61.00 Matches: 11  
Percent Similarity: 72.22% Conservative: 2  
Best Local Similarity: 61.11% Mismatches: 5  
Query Match: 17.48% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1 (1-235) x AD1654 (1-399)

OY 169 AAGACCGCCGATACGCGCATTGGCCGTGACGACGCGCGACTTCACCTGCGAG 222  
DB 368 GlnThrAlaAlaIatYrGlyHisPheGlyArgSerAspLeuAspLeuProTrpGlu 385

RESULT 35

AH1282

S: methionine adenosyltransferase homolog metK [imported] - Listeria monocytogenes (st  
C: Species: Listeria monocytogenes  
C: Date: 27-Nov-2001 #sequence\_revision 27-Nov-2001 #text\_change 14-Dec-2001  
C: Accession: AH1282  
R: Glaaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloec  
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihl,  
D.; Jones, L.M.; Karst, U.  
Science 294, 849-852, 2001  
A: Authors: Kreft, J.; Kuhn, M.; Kunst, F.; Kurapkai, G.; Madueno, E.; Maitournam, A.;  
Ok, C.; Schlueter, T.; Sinoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehla  
A: Title: Comparative genomics of Listeria species.  
A: Reference number: AB1077; MUID: 21537279; PMID: 11679669  
A: Accession: AH1282  
A: Status: preliminary





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Db      367  ArgpHeYrGlnAarPvaLlAlaLAtyGtClYnIshHeGtYlAaGnaAarPLeuAaPLeuPro 386
OY      217  TGGCAG 222
Db      387  TrpGlu 388
|||
RESULT 41
E82319
S:adenosylmethionine synthase VC0472 [imported] - Vibrio cholerae (strain N16961 sero
C:Species: Vibrio cholerae
C:Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 02-Feb-2001
C:Accession: E82319
R:Heidelberg, J.F.; Eisen, J.A.; Nelson, W.C.; Clayton, R.A.; Gwinn, M.L.; Dodson, R.;
Chardson, D.; Ermolaeva, M.D.; Vamathevan, J.; Bass, S.; Qin, H.; Dragoi, I.; Sellers
1, R.R.; Mekalanos, J.J.; Venter, J.C.; Fraser, C.M.
A:Title: DNA Sequence of both chromosomes of the cholera pathogen Vibrio cholerae.
A:Reference number: A82035; MUID:20406833; PMID:10952301
A:Accession: E82319
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-385 <HEI>
A:Cross-references: GB:AE004133; GB:AE003852; MID:9654889; PIDN:AAF93645.1; GSPDB:GN
A:Experimental source: serogroup O1; strain N16961; biotype El Tor
C:Genetics:
A:Gene: VC0472
A:Map position: 1
C:Superfamily: methionine adenosyltransferase

Alignment Scores:
Pred. No.:      3.45      Length:      385
Score:          60.00     Matches:      10
Percent Similarity: 100.00%  Conservative: 2
Best Local Similarity: 83.33%  Mismatches:  0
Query Match:     17.19%     Indels:       0
DB:              2         Gaps:         0

US-09-198-779B-1 (1-235) x E82319 (1-385)

OY      169  AAGACCGCCGCATACGGCCACTTTGGCCGTGACGAC 204
|||
Db      355  LysThrAlaAlaTyrGtYnIshHeGtYlAaGnaAarPLeu 366
|||
RESULT 42
E84381
S:acylaminoacyl-peptidase [imported] - Halobacterium sp. NRC-1
C:Species: Halobacterium sp. NRC-1
C:Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001
C:Accession: E84381
R:Ng, W.V.; Kennedy, S.P.; Mahlairs, G.G.; Berquist, B.; Pan, M.; Shukla, H.D.; Lasky
1; Lelhausser, B.; Keller, K.; Cruz, R.; Danson, M.J.; Hough, D.W.; Maddocks, D.G.; Ja
Jung, K.H.; Alam, M.; Freitas, T.
C:Proc. Natl. Acad. Sci. U.S.A. 97, 12176-12181, 2000
A:Authors: Hou, S.; Daniels, C.J.; Dennis, P.P.; Omer, A.D.; Ehardt, H.; Lowe, T.M.;
A:Title: Genome sequence of Halobacterium species NRC-1.
A:Reference number: A84160; MUID:20504483; PMID:11016950
A:Accession: E84381
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-674 <STO>
A:Cross-references: GB:AE004437; MID:910581715; PIDN:AAG20414.1; GSPDB:GN00138
C:Genetics:
A:Gene: yuxL

Alignment Scores:
Pred. No.:      3.54      Length:      674
Score:          60.00     Matches:      25
Percent Similarity: 31.03%  Conservative: 2
Best Local Similarity: 28.74%  Mismatches:  28
Query Match:     17.34%     Indels:       32
DB:              2         Gaps:         3

US-09-198-779B-1 (1-235) x E84381 (1-674)

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Oy 228 GACCACCTGCGAGTGAAGTCGGCGTCACGCGCAAGTGGCCGTATGCGGC----- 175
    ||| ||||| ||| ||||| ||||| ||||| |||||
Db 183 AspleuAlaAlaGlyArgValAspArgValThrAlaGlyValAlaGlyCysGlyPro 202
Oy 174 -----GCTCTTCAT----- 166
Db 203 AlaTrpGlyAspAspGlyThrLeuTyTrpProIleArgGlyLeuAspAlaasp 222
Oy 165 -----GAACCTGNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNAACGTAAGTGG 112
    ||| ||| ||| ||| |||
Db 223 ArgLeuGluTrpAlaIleGluAlaAlaThrProAlaAspSerAlaAspSerThrValAl 242
Oy 111 TTGTGCTGCTGCTCAATATACACATTAATAACACACAAATATGACGACGCTGC 52
    ||||| ||| ||||| |||
Db 243 -----ThrThrValGluGlyMetClyProthr 251

Oy 51 ATTATATCAGCATGGGACGCT 31
    :: ||||| |||
Db 252 LeuAlaValHisGlySerArg 258

RESULT 43
A71281
probable S-adenosylmethionine synthetase (metk) - syphilis spirochete
C:Species: Treponema pallidum subsp. pallidum (syphilis spirochete)
C:Date: 24-Jul-1998 #sequence_revision 24-Jul-1998 #text_change 18-Jun-1999
C:Accession: A71281
R:Fraser, C.M.; Norris, S.J.; Weinstein, G.M.; White, O.; Sutton, G.G.; Dodson, R.; Gwitrson, J.; Khalak, H.; Richardson, D.; Howell, J.K.; Chidambaram, M.; Utterback, T.; McDCChey, L.; Weidman, J.; Smith, H.O.; Venter, J.C.
Science 281, 375-388, 1998
A:Title: Complete genome sequence of Treponema pallidum, the syphilis spirochete.
A:Reference number: A71250; MUID:98332770; PMID:9665876
C:Accession: A71281
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-396 <COL>
A:Cross-references: GB:AE001250; GB:AE000520; NID:93323096; PIDN:AC65758.1; PID:9332310
A:Experimental source: strain Nichols
C:Genetics:
A:Gene: TP0794
C:Superfamily: methionine adenosyltransferase

Alignment Scores:
Pred. No.: 4.89 Length: 396
Score: 59.00 Matches: 13
Percent Similarity: 54.55% Conservative: 5
Best Local Similarity: 39.39% Mismatches: 7
Query Match: 16.91% Indels: 8
DB: 2 Gaps: 2

US-09-198-779b-1 (1-235) x A71281 (1-396)
Oy 160 AGGTCATCAAGACCGCGCATACGGCATTGGCCGTGACGAC----- 204
    ||||| ||||| ||||| ||||| ||||| |||||
Db 355 ArgTyrArgSerThrAlaValTyrGlyHisPheGlyArgGlnPheProTrpGluArg 374
Oy 205 GCCGACTTCACCTCGGAGGTG-----GTCAGAGCC 234
    ||| ||||| ||||| ||||| |||||
Db 375 ThrAspCysValCysAspLeuGlnArgAlaValArgPro 387

RESULT 44
A47151
methionine adenosyltransferase (EC 2.5.1.6) - mouse
N:Alternate names: S-adenosylmethionine synthetase
C:Species: Mus musculus (house mouse)
C:Date: 21-Jan-1994 #sequence_revision 18-Nov-1994 #text_change 05-May-2000
C:Accession: A47151
R:Sakata, S.F.; Shelly, L.L.; Ruppert, S.; Schutz, G.; Chou, J.Y.
J. Biol. Chem. 268, 13978-13986, 1993
A:Title: Cloning and expression of murine S-adenosylmethionine synthetase.
A:Reference number: A47151; MUID:93300783; PMID:8314764
A:Accession: A47151

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A:Status: preliminary
A:Molecule type: nucleic acid
A:Residues: 1-396 <SAK>
A:Experimental source: liver
A>Note: sequence inconsistent with the nucleotide translation
A>Note: sequence extracted from NCBI backbone (NCBI:134412, NCBI:134413)
C:Superfamily: methionine adenosyltransferase
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:
Pred. No.: 4.89 Length: 396
Score: 59.00 Matches: 13
Percent Similarity: 71.43% Conservative: 2
Best Local Similarity: 61.90% Mismatches: 4
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US-09-198-779b-1 (1-235) x A47151 (1-396)
Oy 169 AAGACGCGCGCATACGCGCATTGGCCGTGACGACGCGGACTTCACCTGCGAGGTGTC 228
    ||||| ||||| ||||| ||||| ||||| |||||
Db 374 LysThrAlaCysTyrGlyHisPheGlyArg-----SerGluPheProTrpGluValPro 391
Oy 229 AAG 231
    |||
Db 392 Lys 392

RESULT 45
S06114
methionine adenosyltransferase (EC 2.5.1.6) - rat
N:Alternate names: S-adenosylmethionine synthetase
C:Species: Rattus norvegicus (Norway rat)
C:Date: 31-Mar-1990 #sequence_revision 31-Mar-1990 #text_change 05-May-2000
C:Accession: S06114; S18256; S18257
R:Hotikawa, S.; Ishikawa, M.; Ozasa, H.; Tsukada, K.
Eur. J. Biochem. 184, 497-501, 1989
A:Title: Isolation of a cDNA encoding the rat liver S-adenosylmethionine synthetase.
A:Reference number: S06114; MUID:90032633; PMID:2806235
C:Accession: S06114
A:Molecule type: mRNA
A:Residues: 1-397 <HOB>
A:Cross-references: EMBL:X15734; NID:957183; PIDN:CA433754.1; PID:957184
R:Mato, J.M.
submitted to the EMBL Data Library, July 1991
A:Reference number: S18256
A:Accession: S18256
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-397 <MAT>
A:Cross-references: EMBL:X60822
R:Alvarez, L.; Asuncion, M.; Corrales, F.; Pajares, M.A.; Mato, J.M.
FEBS Lett. 290, 142-146, 1991
A:Title: Analysis of the 5' non-coding region of rat liver S-adenosylmethionine synth
A:Reference number: S18257; MUID:92008649; PMID:1915866
C:Accession: S18257
A:Status: translation not shown
A:Molecule type: mRNA
A:Residues: 1-46 <ALV>
A:Cross-references: EMBL:X60822
C:Superfamily: methionine adenosyltransferase
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:
Pred. No.: 4.89 Length: 397
Score: 59.00 Matches: 13
Percent Similarity: 71.43% Conservative: 2
Best Local Similarity: 61.90% Mismatches: 4
Query Match: 16.91% Indels: 2
DB: 2 Gaps: 1

US-09-198-779b-1 (1-235) x S06114 (1-397)
Oy 169 AAGACGCGCGCATACGCGCATTGGCCGTGACGACGCGGACTTCACCTGCGAGGTGTC 228

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Db 375 LysThrAlaCysTyrGlyHisPheGlyArg-----SerGluPheProTrpGluValPro 392
QY 229 AAG 231
Db 393 Lys 393
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Search completed: April 23, 2003, 12:07:53  
Job time : 24.5833 secs



PR 29-MAR-1999; 9905-0126785.  
PR 01-APR-1999; 9905-0127462.  
PR 06-APR-1999; 9905-0128234.  
PR 08-APR-1999; 9905-0128714.  
PR 16-APR-1999; 9905-0129845.  
PR 19-APR-1999; 9905-0130077.  
PR 21-APR-1999; 9905-0130449.  
PR 23-APR-1999; 9905-0130510.  
PR 23-APR-1999; 9905-0130891.  
PR 28-APR-1999; 9905-0131449.  
PR 30-APR-1999; 9905-0132048.  
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PR 18-JUN-1999; 9905-0139461.  
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PR 12-JUL-1999; 9905-0142977.  
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PR 14-JUL-1999; 9905-0143624.  
PR 15-JUL-1999; 9905-0144005.  
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PR 16-JUL-1999; 9905-0144086.

PR 19-JUL-1999; 9905-0144325.  
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PR 26-OCT-1999; 99US-0161360.
PR 26-OCT-1999; 99US-0161361.
PR 28-OCT-1999; 99US-0161920.
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PR 28-OCT-1999; 99US-0161993.
PR 29-OCT-1999; 99US-0162142.

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Query Match 97.9%; Score 74.4; DB:21; Length 1674;
Best Local Similarity 98.7%; Pred. No. 5.5e-14;
Matches 75; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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```

OY 1 AGGTTTCATCAGACCGCCGCGCATACGCGCACTTGGCCGTGAGAGAGCCGCGACTTCACCTGC 60
    |||||
DB 1188 AGGTTTCATCAGACCGCCGCGCATACGCGCACTTGGCCGTGAGAGAGCCGCGACTTCACCTGC 1247
    |||||
OY 61 GAGGTGTCAGAGCCCC 76
    |||||
DB 1248 GAGGTGTCAGAGCCCC 1263

```

```

RESULT 2
AA07185 standard; cDNA; 1380 BP.
ID AA07185
XX
AC AA07185;
XX
DT 21-MAY-1999 (first entry)
XX
DE Wheat S-adenosylmethionine synthetase cDNA contig.
XX
KW S-adenosylmethionine synthetase; wheat; amino acid; lysine;
KW threonine; methionine; cysteine; isoleucine; transgenic plant;
KW crop improvement; food; feedstuff; ss.
XX
OS Triticum aestivum.
XX
FH Key Location/Qualifiers
FT CDS 73..1257 /*tag= a
FT
XX
PN MO9855601-A2.
XX
PD 10-DEC-1998.
XX
PF 05-JUN-1998; 98WO-US11692.
XX
PR 12-JUN-1997; 97US-0049443.
PR 06-JUN-1997; 97US-0048771.
XX
PA (DUPO ) DU PONT DE NEMOURS & CO E I.

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```

XX Abell LM, Allen SM, Falco SC, Hitz WD, Kinney AJ;
PI Rafalski JA, Thorpe CJ;
XX WPI: 1999-070263/06.
DR P-PSDB: AAW97744.
XX
XX New plant amino acid biosynthetic enzymes, DNA and chimeric genes -
PT encode: dihydrodipicolinate reductase; diaminopimelate epimerase;
PT threonine synthase; threonine deaminase; S-adenosylmethionine
PT synthetase
XX
PS Claim 47; Page 69-70; 98pp; English.
XX

```

This is the nucleotide sequence of a cDNA contig that codes for a full-length wheat S-adenosylmethionine synthetase (see AAW97744). The contig was assembled from clones isolated from wheat kernel, leaf, seedling and root cDNA libraries and identified by comparison to public sequence databases using BLAST algorithms. It shows sequence similarity to the barley enzyme. The invention relates to new isolated nucleic acid fragments (see AA07168-85) encoding plant enzymes (see AAW97727-44) that catalyze steps in the biosynthesis of lysine, threonine, methionine, cysteine and isoleucine from aspartate, the enzyme being selected from dihydrodipicolinate reductase, diaminopimelate epimerase, threonine synthase, threonine deaminase or S-adenosylmethionine synthetase. The invention also relates to the construction of a chimeric gene encoding all or a portion of the biosynthetic pathway enzyme, in sense or antisense orientation, where expression of the chimeric gene results in production of altered levels of the enzyme in a transformed host cell.

Overexpression or reduction of expression of genes encoding the amino acid biosynthetic pathway enzymes in crop plants such as corn, soybean and wheat can be used to alter levels of the amino acids in human food and animal feed. Transformed host cells can also be used to identify compounds that inhibit one of the enzymes.

```

SQ Sequence 1380 BP; 299 A; 430 C; 379 G; 267 T; 5 other;
Query Match 85.3%; Score 64.8; DB:20; Length 1380;
Best Local Similarity 90.8%; Pred. No. 5.2e-11;
Matches 69; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

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OY 1 AGGTTTCATCAGACCGCCGCGCATACGCGCACTTGGCCGTGAGAGAGCCGCGACTTCACCTGC 60
    |||||
DB 1156 AGGTTTCATCAGACCGCCGCGCATACGCGCACTTGGCCGTGAGAGAGCCGCGACTTCACCTGC 1215
    |||||
OY 61 GAGGTGTCAGAGCCCC 76
    |||||
DB 1216 GAGGTGTCAGAGCCCC 1231

```

```

RESULT 3
AAT99143 standard; cDNA to mRNA; 1182 BP.
ID AAT99143
XX
AC AAT99143;
XX
DT 26-MAR-1998 (first entry)
XX
DE S-adenosylmethionine synthase 3 gene.
XX
KW S-adenosylmethionine synthase 3 gene; barley; alkali resistant plant;
KW sam3; ss.
XX
OS Hordeum vulgare.
XX
PN JP09313186-A.
XX
PD 09-DEC-1997.
XX
PF 28-MAY-1996; 96JP-0133406.
XX
PR 28-MAY-1996; 96JP-0133406.
XX

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XX (NIOC ) NIPPON OIL CO LTD.  
 XX WPI: 1998-080077/08.  
 DR P-PSDB: AAW34542.  
 XX S-adenosyl-methionine synthase gene - useful in producing plants  
 PT resistant to alkaline soil  
 XX Claim 6; Page 10-11; 13pp; Japanese.  
 CC This sequence represents the S-adenosylmethionine synthase 3 (sam3)  
 CC gene. This DNA sequence may be used in a vector to produce plants which  
 CC are resistant to alkaline soil.  
 XX  
 SQ Sequence 1182 BP; 253 A; 368 C; 331 G; 230 T; 0 other;

Query Match 81.1%; Score 61.6; DB 19; Length 1182;  
 Best Local Similarity 86.2%; Pred. No. 5.1e-10;  
 Matches 67; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

QY 1 AGGTTTCATCAGACCGCCGCAATACGCGCACTTTGGCCGTGACGACGCCGACCTTCACCTGC 60  
 |||||  
 DB 1084 AGGTTTCATCAGACGACGCTGCTTATGCTCACTTTGGCCGCGAGCATGCCACTTCACCTGG 1143  
 |||||  
 QY 61 GAGGTGTCAGACCCC 76  
 |||||  
 DB 1144 GAGGTGTCAGACCCC 1159

RESULT 4  
 AAT99141  
 ID AAT99141 standard; cDNA to mRNA; 1182 BP.  
 XX  
 AC AAT99141;  
 XX  
 DT 26-MAR-1998 (first entry)  
 XX  
 DE S-adenosylmethionine synthase 1 gene.  
 XX  
 KW S-adenosylmethionine synthase 1 gene; barley; alkali resistant plant;  
 KW sam1; ss.  
 XX  
 OS Hordeum vulgare.  
 XX  
 PN JP09313186-A.  
 XX  
 PD 09-DEC-1997.  
 XX  
 PF 28-MAY-1996; 96JP-0133406.  
 XX  
 PR 28-MAY-1996; 96JP-0133406.  
 XX  
 PA (NIOC ) NIPPON OIL CO LTD.  
 XX  
 DR WPI: 1998-080077/08.  
 DR P-PSDB: AAW34540.  
 XX  
 PT S-adenosyl-methionine synthase gene - useful in producing plants  
 PT resistant to alkaline soil  
 XX  
 PS Claim 4; Page 8-9; 13pp; Japanese.  
 XX  
 CC This sequence represents the S-adenosylmethionine synthase 1 (sam1)  
 CC gene. This DNA sequence may be used in a vector to produce plants which  
 CC are resistant to alkaline soil.  
 XX  
 SQ Sequence 1182 BP; 253 A; 372 C; 324 G; 233 T; 0 other;

Query Match 78.9%; Score 60; DB 19; Length 1182;  
 Best Local Similarity 86.8%; Pred. No. 1.6e-09;  
 Matches 66; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

QY 1 AGGTTTCATCAGACCGCCGCAATACGCGCACTTTGGCCGTGACGACGCCGACCTTCACCTGC 60  
 |||||  
 DB 1084 AGGTTTCATCAGACGACGCTGCTTATGCTCACTTTGGCCGCGAGCATGCCACTTCACCTGG 1143  
 |||||  
 QY 61 GAGGTGTCAGACCCC 76  
 |||||  
 DB 1144 GAGGTGTCAGACCCC 1159

RESULT 5  
 AAT99142  
 ID AAT99142 standard; cDNA to mRNA; 1182 BP.  
 XX  
 AC AAT99142;  
 XX  
 DT 26-MAR-1998 (first entry)  
 XX  
 DE S-adenosylmethionine synthase 2 gene.  
 XX  
 KW S-adenosylmethionine synthase 2 gene; barley; alkali resistant plant;  
 KW sam2; ss.  
 XX  
 OS Hordeum vulgare.  
 XX  
 PN JP09313186-A.  
 XX  
 PD 09-DEC-1997.  
 XX  
 PF 28-MAY-1996; 96JP-0133406.  
 XX  
 PR 28-MAY-1996; 96JP-0133406.  
 XX  
 PA (NIOC ) NIPPON OIL CO LTD.  
 XX  
 DR WPI: 1998-080077/08.  
 DR P-PSDB: AAW34541.  
 XX  
 PT S-adenosyl-methionine synthase gene - useful in producing plants  
 PT resistant to alkaline soil  
 XX  
 PS Claim 5; Page 9-10; 13pp; Japanese.  
 XX  
 CC This sequence represents the S-adenosylmethionine synthase 2 (sam2)  
 CC gene. This DNA sequence may be used in a vector to produce plants which  
 CC are resistant to alkaline soil.  
 XX  
 SQ Sequence 1182 BP; 261 A; 368 C; 324 G; 229 T; 0 other;

Query Match 76.8%; Score 58.4; DB 19; Length 1182;  
 Best Local Similarity 85.5%; Pred. No. 5e-09; Indels 11; Gaps 0;  
 Matches 65; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

QY 1 AGGTTTCATCAGACCGCCGCAATACGCGCACTTTGGCCGTGACGACGCCGACCTTCACCTGC 60  
 |||||  
 DB 1084 AGGTTTCATCAGACGACGCTGCTTATGCTCACTTTGGCCGCGAGCATGCCACTTCACCTGG 1143  
 |||||  
 QY 61 GAGGTGTCAGACCCC 76  
 |||||  
 DB 1144 GAGGTGTCAGACCCC 1159

RESULT 6  
 AAX81128  
 ID AAX81128 standard; cDNA; 399 BP.  
 XX  
 AC AAX81128;  
 XX  
 DT 06-SEP-1999 (first entry)  
 XX  
 DE S-adenosyl methionine (SAM) partial cDNA sequence.  
 XX  
 KW Senescence-associated receptor-like protein kinase; sakr; sam gene; sag;  
 KW senescence-associated gene; plant senescence; promoter; pharmaceutical;

Query Match 78.9%; Score 60; DB 19; Length 1182;  
 Best Local Similarity 86.8%; Pred. No. 1.6e-09;  
 Matches 66; Conservative 0; Mismatches 10; Indels 0; Gaps 0;



KW plant maturation; S-adenosyl methionine; flower; fruit development; ss.  
 OS Phaseolus vulgaris.  
 XX MO9929159-A1.  
 XX 17-JUN-1999.  
 XX PD  
 XX PF 08-DEC-1998; 98WO-US25799.  
 XX PR 08-DEC-1997; 97US-0067898.  
 XX (VITA-) VITALITY BIOTECHNOLOGIES INC.  
 PA Gepstein S, Hajuoje T, Rosner A;  
 PI WPI: 1999-404873/34.  
 DR P-PSDB; AAY21978.  
 XX  
 PT DNA encoding senescence-associated genes for a senescence  
 PT receptor-like protein kinase  
 XX  
 PS Claim 18; Fig 4; 70pp; English.  
 XX The invention relates to a senescence-associated receptor-like protein  
 CC kinase (sark) gene. The sark gene is a senescence-associated gene (sag)  
 CC and is expressed early in the plant senescence process. The sark gene  
 CC promoter is useful for driving expression of foreign genes having a  
 CC desired product, such as a pharmaceutical, during the process of plant  
 CC maturation. The sark gene promoter can be used to drive expression of  
 CC resistance genes against pathogens or pests during senescence when the  
 CC plant is particularly susceptible to infection or infestation. The sark  
 CC gene promoter may also be used to drive expression of a gene encoding an  
 CC inhibitor of senescence. Plant senescence may be inhibited by use of  
 CC antisense sark constructs. Over expression of the sag genes, using the  
 CC sark or sam (S-adenosyl methionine) gene promoters is useful for  
 CC induction of early senescence. This is useful to obtain flower or fruit  
 CC development prior to specific pest onset, prior to undesirable cross-  
 CC fertilization from related crops, at a specific time during storage or  
 CC retail, or to avoid development of plant structures that are not of  
 CC agronomic importance. The present sequence represents a partial cDNA  
 CC clone of S-adenosyl methionine (SAM).  
 XX  
 SO Sequence 399 BP; 108 A; 80 C; 82 G; 129 T; 0 other;  
 Query Match 70.5%; Score 53.6; DB 20; Length 399;  
 Best Local Similarity 81.6%; Pred. No. 1.4e-07;  
 Matches 62; Conservative 0; Mismatches 14; Indels 0; Gaps 0;  
 QY 1 AGTTCATCAAGACCGCCGATAGCGCACTTGGCCGTGACGAGCGCCGACTTCACCTGC 60  
 DB 102 AGTTCCTTGAGACGCTGCATATGACACCTTCGCGACAGAGGAGCGACTTCACATGG 161  
 QY 61 GAGGTGCTCAAGCCCC 76  
 DB 162 GAAGTGTCAAGCCCC 177  
 RESULT 7  
 ID AAX07184 standard; cDNA; 1485 BP.  
 XX AAX07184;  
 XX 21-MAY-1999 (first entry)  
 XX Soybean S-adenosylmethionine synthetase cDNA clone s2.12b06.  
 DE S-adenosylmethionine synthetase; soybean; amino acid; lysine;  
 KW threonine; methionine; cysteine; isoleucine; transgenic plant;  
 KW crop improvement; food; feedstuff; ss.  
 XX Glycine max.  
 OS

XX Key Location/Qualifiers  
 FH CDS 74..1252  
 FT /\*\*tag- a  
 XX MO9855601-A2.  
 XX 10-DEC-1998.  
 XX PD  
 XX PF 05-JUN-1998; 98WO-US11692.  
 XX PR 12-JUN-1997; 97US-0049443.  
 XX PR 06-JUN-1997; 97US-0048771.  
 XX (DUPO ) DU PONT DE NEMOURS & CO E. I.  
 PA Abell LM, Allen SM, Falco SC, Hiltz WD, Kinney AJ;  
 PI Rafalski JA, Thorpe CJ;  
 DR WPI: 1999-070263/06.  
 DR P-PSDB; AAW97743.  
 XX  
 PT New plant amino acid biosynthetic enzymes, DNA and chimeric genes -  
 PT encode: dihydropicolinate reductase; diaminopimelate epimerase;  
 PT threonine synthase; threonine deaminase; S-adenosylmethionine  
 PT synthetase  
 XX  
 PS Claim 44; Page 66-67; 98pp; English.  
 XX This is the nucleotide sequence of cDNA clone s2.12b06, which  
 CC codes for a full-length soybean S-adenosylmethionine synthetase  
 CC (see AAW97743). The clone was isolated from a soybean seed cDNA  
 CC library and identified by comparison to public sequence databases  
 CC using BLAST algorithms. It shows sequence similarity to the  
 CC tomato enzyme. The invention relates to new isolated nucleic  
 CC acid fragments (see AAX07168-85) encoding plant enzymes (see  
 CC AAW97727-44) that catalyze steps in the biosynthesis of lysine,  
 CC threonine, methionine, cysteine and isoleucine from aspartate, the  
 CC enzyme being selected from dihydropicolinate reductase,  
 CC diaminopimelate epimerase, threonine synthase, threonine deaminase  
 CC or S-adenosylmethionine synthetase. The invention also relates to  
 CC the construction of a chimeric gene encoding all or a portion of  
 CC the biosynthetic pathway enzyme, in sense or antisense orientation,  
 CC where expression of the chimeric gene results in production of  
 CC altered levels of the enzyme in a transformed host cell.  
 CC Overexpression or reduction of expression of genes encoding the  
 CC amino acid biosynthetic pathway enzymes in crop plants such as  
 CC corn, soybean and wheat can be used to alter levels of the amino  
 CC acids in human food and animal feed. Transformed host cells can  
 CC also be used to identify compounds that inhibit one of the enzymes.  
 XX  
 SO Sequence 1485 BP; 366 A; 373 C; 357 G; 389 T; 0 other;  
 Query Match 68.4%; Score 52; DB 20; Length 1485;  
 Best Local Similarity 80.3%; Pred. No. 5.1e-07;  
 Matches 61; Conservative 0; Mismatches 15; Indels 0; Gaps 0;  
 QY 1 AGTTCATCAAGACCGCCGATAGCGCACTTGGCCGTGACGAGCGCCGACTTCACCTGC 60  
 DB 1157 AGTTCCTTGAGACGCTGCATATGACACCTTCGCGACAGAGGAGCGACTTCACATGG 1216  
 QY 61 GAGGTGCTCAAGCCCC 76  
 DB 1217 GAAGTGTCAAGCCCC 1232  
 RESULT 8  
 ID AAA51037 standard; cDNA; 1518 BP.  
 XX AAA51037;  
 AC 09-OCT-2000 (first entry)  
 XX

XX Soybean S-adenosyl-L-methionine synthetase cDNA.  
 DE S-adenosyl-L-methionine synthetase; SAMS; probe; promoter; embryo;  
 KW constitutive; tissue-specific; development-specific;  
 KM herbicide resistance; pathogen resistance; ss.  
 XX Glycine max.  
 OS  
 XX  
 FH Key Location/Qualifiers  
 FT 5'UTR 1..73  
 FT /\*tag= a  
 FT CDS 74..1252  
 FT /\*tag= b  
 FT /product= S-adenosyl-L-methionine\_synthetase  
 PN WO200037662-A2.  
 XX  
 XX 29-JUN-2000.  
 XX  
 XX 17-DEC-1999; 99WO-US30180.  
 XX  
 XX 21-DEC-1998; 98US-0113045.  
 XX  
 XX (DUPO ) DU PONT DE NEMOURS & CO E I.  
 XX  
 XX Falco SC, Li Z;  
 PI  
 DR WPI: 2000-442682/38.  
 XX  
 XX S-adenosyl-L-methionine synthetase promoter for expressing target  
 PT heterologous herbicide-resistance or pathogen-resistance nucleic acid  
 PT fragments in plants, especially soybean  
 XX  
 XX Example 2: Page 39; 50pp: English.  
 PS  
 XX This is the soybean full-length S-adenosyl-L-methionine synthetase (SAMS)  
 CC cDNA, which was used to generate a probe to isolate a SAMS promoter. The  
 CC SAMS promoter is active in seedlings and callus and over-expression of a  
 CC gene in embryo stage can be achieved at an early developing stage using  
 CC the SAMS promoter. The SAMS promoter may be used as an alternative to  
 CC cauliflower mosaic virus 35S promoter to drive expression of selectable  
 CC marker genes. Plant cells transformed with the SAMS constitutive promoter  
 CC are useful for increasing or decreasing the expression of heterologous  
 CC nucleic acid fragments in a plant, preferably corn, rice, wheat, barley,  
 CC palm, Arabidopsis, soybean, oil seed Brassica, peanut, sunflower,  
 CC safflower, cotton, tobacco, tomato, potato or cocoa. Target heterologous  
 CC nucleic acid fragments include herbicide or pathogen resistance  
 CC nucleic acid fragments.  
 XX  
 SQ Sequence 1518 BP; 399 A; 373 C; 357 G; 389 T; 0 other;  
 Query Match 68.4%; Score 52; DB 21; Length 1518;  
 Best Local Similarity 80.3%; Pred. No. 5.1e-07;  
 Matches 61; Conservative 0; Mismatches 15; Indels 0; Gaps 0;  
 QY 1 AGGTCATCAAGACCGCCGACATGCGCCCTTGGCCGTCAGACGCCGACTTCACCTGC 60  
 DB 1157 AGGTCATCAAGACCGCCGACATGCGCCCTTGGCCGTCAGACGCCGACTTCACCTGC 60  
 QY 61 GAGGTGTCAGCCCC 76  
 DB 1217 GAGGTGTCAGCCCC 1232

RESULT 9  
 AAX07183  
 ID AAX07183 standard; cDNA; 1582 BP.  
 XX  
 AC AAX07183;  
 XX  
 DT 21-MAY-1999 (first entry)  
 XX

DE Corn S-adenosylmethionine synthetase cDNA clone cc3.mn0002.d2.  
 XX  
 KW S-adenosylmethionine synthetase; corn; maize; amino acid; lysine;  
 KM threonine; methionine; cysteine; isoleucine; transgenic plant;  
 KM crop improvement; food; feedstuff; ss.  
 XX  
 OS Zea mays.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 140..1330  
 FT /\*tag= a  
 FT  
 PN W09855601-A2.  
 XX  
 XX 10-DEC-1998.  
 XX  
 XX 05-JUN-1998; 98WO-US11692.  
 XX  
 XX 12-JUN-1997; 97US-0049443.  
 XX  
 XX 06-JUN-1997; 97US-0048771.  
 XX  
 XX (DUPO ) DU PONT DE NEMOURS & CO E I.  
 XX  
 XX Abell LM, Allen SM, Falco SC, Hitz WD, Kinney AJ;  
 PI Ratalski JA, Thorpe CJ;  
 DR WPI: 1999-070263/06.  
 DR P-PSDB; AAW97742.  
 XX  
 XX New plant amino acid biosynthetic enzymes, DNA and chimeric genes -  
 PT encode: dihydripycolinate reductase; diaminopimelate epimerase;  
 PT threonine synthase; threonine deaminase; S-adenosylmethionine  
 PT synthetase  
 XX  
 XX Claim 41; Page 62-63; 98pp: English.  
 PS  
 XX This is the nucleotide sequence of cDNA clone cc3.mn0002.d2, which  
 CC codes for a full-length corn S-adenosylmethionine synthetase  
 CC (see AAW97742). The clone was isolated from a corn callus cDNA  
 CC library and identified by comparison to public sequence databases  
 CC using BLAST algorithms. It shows sequence similarity to the  
 CC Oryza sativa enzyme. The invention relates to new isolated  
 CC nucleic acid fragments (see AAX07168-85) encoding plant enzymes (see  
 CC AAW97727-44) that catalyze steps in the biosynthesis of lysine,  
 CC threonine, methionine, cysteine and isoleucine from aspartate, the  
 CC enzyme being selected from dihydripycolinate reductase,  
 CC diaminopimelate epimerase, threonine synthase, threonine deaminase  
 CC or S-adenosylmethionine synthetase. The invention also relates to  
 CC the construction of a chimeric gene encoding all or a portion of  
 CC the biosynthetic pathway enzyme, in sense or antisense orientation,  
 CC where expression of the chimeric gene results in production of  
 CC altered levels of the enzyme in a transformed host cell.  
 CC Overexpression or reduction of expression of genes encoding the  
 CC amino acid biosynthetic pathway enzymes in crop plants such as  
 CC corn, soybean and wheat can be used to alter levels of the amino  
 CC acids in human food and animal feed. Transformed host cells can  
 CC also be used to identify compounds that inhibit one of the enzymes.  
 XX  
 SQ Sequence 1582 BP; 340 A; 474 C; 417 G; 351 T; 0 other;  
 Query Match 67.1%; Score 51; DB 20; Length 1582;  
 Best Local Similarity 80.0%; Pred. No. 1e-06;  
 Matches 60; Conservative 0; Mismatches 15; Indels 0; Gaps 0;  
 QY 2 GGTTCATCAAGACCGCCGACATGCGCCCTTGGCCGTCAGACGCCGACTTCACCTGC 61  
 DB 1230 GGTTCATCAAGACCGCCGACATGCGCCCTTGGCCGTCAGACGCCGACTTCACCTGC 61  
 QY 62 AGGTGTCAGCCCC 76  
 DB 1290 AGGTGTCAGCCCC 1304

RESULT 10  
AAC44219  
ID AAC44219 standard; DNA: 635 BP.  
XX  
AC AAC44219;  
XX  
DT 18-OCT-2000 (first entry)  
XX  
DE Arabidopsis thaliana DNA fragment SEQ ID NO: 42062.  
XX  
XX Hybridisation assay; genetic mapping; gene expression control;  
KW protein identification; signal transduction pathway;  
KW metabolic pathway; promoter; termination sequence; ss.  
XX  
OS Arabidopsis thaliana.  
XX  
PN EPI033405-A2.  
XX  
PD 06-SEP-2000.  
XX  
PF 25-FEB-2000; 2000EP-0301439.  
XX  
PR 25-FEB-1999; 99US-0121825.  
PR 05-MAR-1999; 99US-0123180.  
PR 09-MAR-1999; 99US-0123548.  
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PR 25-MAR-1999; 99US-0126264.  
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PR 01-APR-1999; 99US-0127462.  
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PR 08-APR-1999; 99US-0128714.  
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PR	29-OCT-1999;	9905-0162142.

XX	AC	AAC33986;	
XX	DT	17-OCT-2000	(first entry)
XX	XX		
DE	XX	Arabidopsis thaliana	DNA fragment SEQ ID NO: 5035.
XX	KM	Hybridisation assay; genetic mapping; gene expression control	
XX	KW	protein identification; signal transduction pathway; ss.	
XX	OS	metabolic pathway; promoter; termination sequence; ss.	
XX	PN	Arabidopsis thaliana.	
XX	PD	EP1033405-A2.	
XX	PF	06-SEP-2000.	
XX	PF	25-FEB-2000;	2000EP-0301439.
PR	XX	25-FEB-1999;	99US-0121825.
PR	XX	05-MAR-1999;	99US-0123180.
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PR	XX	28-MAY-1999;	99US-0136782.
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PR 12-JUL-1999; 99US-0142977.  
PR 13-JUL-1999; 99US-0143542.  
PR 14-JUL-1999; 99US-0143624.  
PR 15-JUL-1999; 99US-0144005.  
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PR 23-JUL-1999; 99US-0145224.  
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PR 02-AUG-1999; 99US-0146388.  
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PR 04-AUG-1999; 99US-0147302.  
PR 05-AUG-1999; 99US-0147192.  
PR 05-AUG-1999; 99US-0147260.  
PR 06-AUG-1999; 99US-0147303.  
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QY 61 GAGGTGTCAGACCCC 76  
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DT 18-OCT-2000 (first entry)  
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XX Hybridisation assay: genetic mapping; gene expression control;  
KM protein identification; signal transduction pathway;  
KW metabolic pathway; promoter; termination sequence; ss.  
XX Arabidopsis thaliana.  
OS  
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XX 06-SEP-2000.  
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DT	17-OCT-2000 (first entry)				
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DE	Arabidopsis thaliana DNA Fragment SEQ ID NO: 3408.				

XX	Hybridisation assay; genetic mapping; gene expression control;
KW	protein identification; signal transduction pathway;
KW	metabolic pathway; promoter; termination sequence; ss.
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PN	Ep1033405-A2.
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PD	06-SEP-2000.
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PR 22-OCT-1999; 99US-0160980.  
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PR 26-OCT-1999; 99US-0161360.  
PR 26-OCT-1999; 99US-0161361.  
PR 28-OCT-1999; 99US-0161920.  
PR 28-OCT-1999; 99US-0161992.  
PR 28-OCT-1999; 99US-0161993.  
PR 29-OCT-1999; 99US-0162142.

Query Match 66.3% Score 50.4; DB 21; Length 1529;  
Best Local Similarity 78.9%; Pred. No. 1.6e-06;  
Matches 60; Conservative 0; Mismatches 16; Indels 0; Gaps 0;

OY 1 AGGTCATCAAGACCGCGCATAGCGGCACCTTGGCGTGACGACGCCGACTTCACCTGC 60  
DB 1210 AGGTCCTTGAACACGCGCGCTTACGACACTTTGGAGAGACGACCCGACTTACCTGG 1269

OY 61 GAGGTGTCAGACCCC 76  
DB 1270 GAGGTGTCAGACCCAC 1285

RESULT 14  
AAC45944  
ID AAC45944 standard; DNA; 1653 BP.

AC AAC45944;

DT 18-OCT-2000 (first entry)

XX Arabidopsis thaliana DNA fragment SEQ ID NO: 48340.

DE Hybridisation assay; genetic mapping; gene expression control;

KW protein identification; signal transduction pathway;



KW metabolic pathway; promoter; termination sequence; ss.  
XX  
OS Arabidopsis thaliana.  
XX  
PN EPI033405-A2.  
XX  
PD 06-SEP-2000.  
XX  
PF 25-FEB-2000; 2000EP-0301439.  
XX  
PR 25-FEB-1999; 99US-0121825.  
PR 05-MAR-1999; 99US-0123160.  
PR 09-MAR-1999; 99US-0123548.  
PR 23-MAR-1999; 99US-0125788.  
PR 25-MAR-1999; 99US-0126264.  
PR 29-MAR-1999; 99US-0126785.  
PR 01-APR-1999; 99US-0127462.  
PR 06-APR-1999; 99US-0128234.  
PR 08-APR-1999; 99US-0128714.  
PR 16-APR-1999; 99US-0129845.  
PR 19-APR-1999; 99US-0130077.  
PR 21-APR-1999; 99US-0130449.  
PR 23-APR-1999; 99US-0130510.  
PR 28-APR-1999; 99US-0130891.  
PR 30-APR-1999; 99US-0131449.  
PR 30-APR-1999; 99US-0132048.  
PR 04-MAY-1999; 99US-0132407.  
PR 05-MAY-1999; 99US-0132484.  
PR 06-MAY-1999; 99US-0132485.  
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PR 06-MAY-1999; 99US-0132487.  
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PR 02-JUL-1999; 99US-0142055.  
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PR 03-AUG-1999; 99US-0147038.  
PR 04-AUG-1999; 99US-0147204.  
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PR 23-AUG-1999; 99US-0149930.  
PR 25-AUG-1999; 99US-0150566.  
PR 26-AUG-1999; 99US-0150884.  
PR 27-AUG-1999; 99US-0151066.  
PR 27-AUG-1999; 99US-0151065.  
PR 27-AUG-1999; 99US-0151080.  
PR 30-AUG-1999; 99US-0151303.  
PR 31-AUG-1999; 99US-0151438.  
PR 01-SEP-1999; 99US-0151930.  
PR 07-SEP-1999; 99US-0152363.

PR 10-SEP-1999; 990S-0153070.  
PR 13-SEP-1999; 990S-0153758.  
PR 15-SEP-1999; 990S-0154018.  
PR 16-SEP-1999; 990S-0154039.  
PR 20-SEP-1999; 990S-0154779.  
PR 22-SEP-1999; 990S-0155139.  
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PR 28-SEP-1999; 990S-0156458.  
PR 29-SEP-1999; 990S-0156596.  
PR 04-OCT-1999; 990S-0157117.  
PR 05-OCT-1999; 990S-0157753.  
PR 06-OCT-1999; 990S-0157863.  
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PR 12-OCT-1999; 990S-0158369.  
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PR 26-OCT-1999; 990S-0161361.  
PR 28-OCT-1999; 990S-0161920.  
PR 28-OCT-1999; 990S-0161992.  
PR 28-OCT-1999; 990S-0161993.  
PR 29-OCT-1999; 990S-0162142.

Query Match 64.2%; Score 48.8; DB 21; Length 1653;  
Best Local Similarity 77.6%; Pred. No. 5,1e-06;  
Matches 59; Conservative 0; Mismatches 17; Indels 0; Gaps 0;

OY 1 AGCTTCATCAACACCGCCGATACGCGCTTGGCGGTGAGACGCGCCTTCACCTGC 60  
DB 1228 AGGTTCCAGAAACCGCTGATGCGCATTCGCGCGATGACCGCTTCACCTTGG 1287  
OY 61 GAGGTGTCACGCCC 76  
DB 1288 GAGGTTGTCAACGCC 1303

RESULT 15  
AAC35348  
ID AAC35348 standard; DNA; 1654 BP.  
XX AAC35348;  
XX 17-OCT-2000 (first entry)  
XX Arabidopsis thaliana DNA fragment SFQ ID NO: 9862.  
XX Hybridisation assay; genetic mapping; gene expression control;  
KW protein identification; signal transduction pathway;  
KW metabolic pathway; promoter; termination sequence; ss.  
XX Arabidopsis thaliana.  
OS

XX EPI033405-A2.  
XX 06-SEP-2000.  
PD 25-FEB-2000; 2000EP-0301439.  
PF 25-FEB-1999; 990S-0121825.  
XX 05-MAR-1999; 990S-0123180.  
PR 09-MAR-1999; 990S-0123548.  
PR 23-MAR-1999; 990S-0125788.  
PR 25-MAR-1999; 990S-0126264.  
PR 29-MAR-1999; 990S-0126785.  
PR 01-APR-1999; 990S-0127462.  
PR 06-APR-1999; 990S-0128234.  
PR 08-APR-1999; 990S-0128714.  
PR 16-APR-1999; 990S-0129845.  
PR 19-APR-1999; 990S-0130077.  
PR 21-APR-1999; 990S-0130449.  
PR 23-APR-1999; 990S-0130510.  
PR 28-APR-1999; 990S-0130891.  
PR 30-APR-1999; 990S-0131449.  
PR 30-APR-1999; 990S-0132048.  
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PR 05-MAY-1999; 990S-0132484.  
PR 06-MAY-1999; 990S-0132485.  
PR 06-MAY-1999; 990S-0132486.  
PR 07-MAY-1999; 990S-0132487.  
PR 11-MAY-1999; 990S-0134286.  
PR 14-MAY-1999; 990S-0134218.  
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PR 27-MAY-1999; 990S-0136392.  
PR 28-MAY-1999; 990S-0136782.  
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PR 03-JUN-1999; 990S-0137528.  
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PR 23-JUN-1999; 990S-0140353.  
PR 23-JUN-1999; 990S-0140354.  
PR 24-JUN-1999; 990S-0140695.  
PR 28-JUN-1999; 990S-0140823.  
PR 29-JUN-1999; 990S-0140991.  
PR 30-JUN-1999; 990S-0141287.

PR 01-JUL-1999; 99US-0141842.  
PR 01-JUL-1999; 99US-0142154.  
PR 02-JUL-1999; 99US-0142055.  
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PR 28-OCT-1999; 99US-0161920.  
PR 28-OCT-1999; 99US-0161920.  
PR 28-OCT-1999; 99US-0161993.  
PR 29-OCT-1999; 99US-0162142.

Query Match Score 64.2%; DB 21; Length 1654;  
Best Local Similarity 77.6%; Pred. No. 5,1e-06;  
Matches 59; Conservative 0; Mismatches 17; Indels 0; Gaps 0;

OY 1 AGGTTCAATCAAGACCGCGCATACGCCACATTGGCCGTGACGACGCCGACTTACCTGC 60  
Db 1229 AGGTTCAAGAAACCGCGCATACGCCACATTGGCGCGTGATGACCCCTTCACTTGG 1288  
OY 61 GAGGTGTCAGGCC 76  
Db 1289 GAGGTGTCAGGCC 1304

RESULT 16  
AAD02296  
AAD02296 standard; DNA; 1636 BP.  
XX AAD02296;  
AC 28-MAR-2001 (first entry)  
DT XX  
XX Nicotiana tabacum S-adenosylmethionine synthetase (SAMS) DNA.  
DE XX  
XX Nicotiana tabacum S-adenosylmethionine synthetase; SAMS; ds.  
KW Tobacco; alkaloid; nicotine; transgenic plant; pharmaceutical protein;  
KW herbicide resistance; S-adenosylmethionine synthetase; SAMS; ds.  
OS Nicotiana tabacum.  
XX  
XX  
FH Key Location/Qualifiers  
FT CDS 96..1268  
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PR	25-MAR-1999	9905-0126764
PR	29-MAR-1999	9905-0126785
PR	01-APR-1999	9905-0127462
PR	06-APR-1999	9905-0128234
PR	08-APR-1999	9905-0128714
PR	16-APR-1999	9905-0129845
PR	19-APR-1999	9905-0130077
PR	21-APR-1999	9905-0130349
PR	23-APR-1999	9905-0130510
PR	28-APR-1999	9905-0130891
PR	28-APR-1999	9905-0131449
PR	30-APR-1999	9905-0132048
PR	30-APR-1999	9905-0132407
PR	04-MAY-1999	9905-0132484
PR	05-MAY-1999	9905-0132485
PR	06-MAY-1999	9905-0132486
PR	07-MAY-1999	9905-0132487
PR	07-MAY-1999	9905-0132487
PR	11-MAY-1999	9905-0134256
PR	14-MAY-1999	9905-0134218
PR	14-MAY-1999	9905-0134219
PR	14-MAY-1999	9905-0134221
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DT 17-OCT-2000 (first entry)
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KW protein identification; signal transduction pathway;
KW metabolic pathway; promoter; termination sequence; ss.
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OS Arabidopsis thaliana.
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PD 06-SEP-2000;
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KW MT; methylene tetrahydrofolate reductase; MTHR; activated methyl cycle;  
KW tylosin production; ds.  
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PR 13-NOV-1996; 96US-030898P.  
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PA (ELIL ) LILLY & CO ELI.  
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PI Dehoff BS, Rosteck PR;  
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XX WPI: 2002-024904/03.  
DR P-PSDB; AAE13583, AAE13584, AAE13585.  
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PT New S-adenosylmethionine (SAM) operon from Streptomyces fradiae which  
PT encodes SAM synthetase, methyltransferase and methylene  
PT tetrahydrofolate reductase, useful for producing SAM by recombinant  
PT techniques  
XX  
PS Claim 19: Column 11-20; 22pp; English.  
XX  
CC The patent discloses Streptomyces fradiae S-adenosylmethionine (SAM)  
CC operon which comprises three genes encoding SAM synthetase, methyl-  
CC transferase (MT) and methylene tetrahydrofolate reductase (MTHR).  
CC SAM synthetase, MT and MTHR together comprise the activated methyl  
CC cycle which produces SAM and provides methyl groups required for  
CC the final steps in tylosin production. The invention also relates  
CC to vectors and transformed heterologous host cells for expressing  
CC SAM synthetase. It also relates to a method useful for producing  
CC SAM by recombinant techniques. The present DNA sequence is SAM  
CC operon from Streptomyces fradiae.  
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SQ Sequence 4848 BP; 648 A; 1869 C; 1696 G; 635 T; 0 other;

OY 4 TTTCATCAAGACCGCCGCTACGCGCCTTTGGCCGTGACGACGCCGACTTCACCTGCGA 62  
DB 2105 TACGCCAAGACCGCCGCTACGCGCCTTCGCGCGCACTGCGGAGTTCACCTGCGGA 2163  
Query Match 50.3%; Score 38.2; DB 24; Length 4848;  
Best Local Similarity 78.0%; Pred. No. 0.011;  
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XX  
DT 26-FEB-2002 (first entry)  
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XX  
KW S-adenosylmethionine; SAM operon; SAM synthetase; methyltransferase;  
KW MT; methylene tetrahydrofolate reductase; MTHR; activated methyl cycle;  
KW tylosin production; ss.  
XX  
OS Streptomyces fradiae.  
XX  
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PN  
PD 06-NOV-2001.  
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PF 22-OCT-1997; 97US-0955957.  
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PR 13-NOV-1996; 96US-030898P.  
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PA (ELIL ) LILLY & CO ELI.  
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PI Dehoff BS, Rosteck PR;  
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XX WPI: 2002-024904/03.

XX New S-adenosylmethionine (SAM) operon from Streptomyces fradiae which  
 PT encodes SAM synthetase, methyltransferase and methylene  
 PT tetrahydrofolate reductase, useful for producing SAM by recombinant  
 PT techniques  
 PS Claim 19; Column 31-36; 22pp; English.  
 XX  
 CC The patent discloses Streptomyces fradiae S-adenosylmethionine (SAM)  
 CC operon which comprises three genes encoding SAM synthetase, methyl-  
 CC transferase (MT) and methylene tetrahydrofolate reductase (MTHFR).  
 CC SAM synthetase, MT and MTHFR together comprise the activated methyl  
 CC cycle which produces SAM and provides methyl groups required for  
 CC the final steps in lysosin production. The invention also relates  
 CC to vectors and transformed heterologous host cells for expressing  
 CC SAM synthetase. It also relates to a method useful for producing  
 CC SAM by recombinant techniques. The present mRNA sequence is SAM  
 CC operon from Streptomyces fradiae.  
 CC  
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 AC AA75637;  
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 DT 22-JAN-2001 (first entry)  
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 KW antibiotic; C12-hydroxylase; pick; desosamine biosynthesis;  
 KW desosaminyl transferase enzyme; ketolide; beta-glucosidase enzyme;  
 KW picromycin biosynthesis; ss.  
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 OS Streptomyces venezuelae.  
 XX  
 PN US6117659-A.  
 PD 12-SEP-2000.  
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 PF 27-MAY-1999; 99US-0320878.  
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 PR 28-MAY-1998; 98US-0087080.  
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 XX  
 PA (KOSA-) KOSAN BIOSCIENCES INC.  
 PI Ashley G, Betlach MC, Betlach M, Tang L, McDaniel R;  
 DR WPI: 2000-610844/58.  
 XX  
 PT New recombinant pick hydroxylase gene of Streptomyces venezuelae useful  
 PT for converting ketolides to antibiotics and as antibiotics and  
 PT intermediates in the synthesis of compounds with pharmaceutical value  
 XX  
 PS Disclosure: Columns 45-46; 117pp; English.

XX The present sequence is used to produce the recombinant DNA compounds  
 CC of the invention. The specification describes a recombinant DNA compound  
 CC expressing recombinant polyketide synthase genes in host cells for the  
 CC production of narbonolide, narbonolide derivatives in the polyketides that  
 CC are useful as antibiotics and as intermediates in the synthesis of  
 CC compounds with pharmaceutical value. The DNA compounds may also encode  
 CC a C12-hydroxylase (pick), desosamine biosynthesis and desosaminyl  
 CC transferase enzymes (useful for conversion of ketolides to antibiotics),  
 CC and the beta-glucosidase enzyme (involved in picromycin biosynthesis).  
 CC These compounds are also useful for increasing the antibiotic activity  
 CC of a compound relative to the unhydroxylated compound. The recombinant  
 CC host cells are useful as genetic systems that allow rapid engineering  
 CC of the narbonolide polyketide synthase. These would be valuable for  
 CC creating novel ketolide analogs for pharmaceutical applications.  
 CC  
 SQ Sequence 1693 BP; 237 A; 566 C; 633 G; 257 T; 0 other;  
 Query Match 49.5%; Score 37.6; DB 21; Length 1693;  
 Best Local Similarity 76.7%; Pred. No. 0.015;  
 Matches 46; Conservative 0; Mismatches 14; Indels 0; Gaps 0;  
 QY 4 TTCATCAGACCGCCGCTAGCGCCACTTGGCGGTACGACGCCGACTTCACCTGGCA 63  
 DB 798 TACTCCAGACCGCCGCTAGCGCCACTTGGCGGTACGACGCCGACTTCACCTGGCA 739  
 RESULT 22  
 AA256005/c  
 ID AA256005 standard; DNA; 1693 BP.  
 AC AA256005;  
 XX  
 DT 23-MAR-2000 (first entry)  
 DE Contig 004 from cosmid PKOS023-27 from Streptomyces venezuelae.  
 XX  
 KW Narbonolide polyketide synthase; PKS; cosmid PKOS023-27; contig 004;  
 KW ketolide; SAM synthase; S-adenosylmethionine synthase; hydroxylase;  
 KW picromycin; antibiotic production; narbonolide; ds.  
 XX  
 OS Streptomyces venezuelae.  
 XX  
 PN W09961599-A2.  
 PD 02-DEC-1999.  
 XX  
 PF 27-MAY-1999; 99WO-US11814.  
 XX  
 PR 28-MAY-1998; 98US-0087080.  
 PR 28-AUG-1998; 98US-0141908.  
 PR 22-SEP-1998; 98US-0100880.  
 PR 08-FEB-1999; 99US-0119139.  
 XX  
 PA (KOSA-) KOSAN BIOSCIENCES INC.  
 PI Ashley G, Betlach MC, Betlach M, McDaniel R, Tang L;  
 DR WPI: 2000-072618/06.  
 DR P-PSDB: AAY67216, AAY67217.  
 XX  
 PT New recombinant DNA encoding a domain of narbonolide polyketide  
 PT synthase, for production of ketolide antibiotics -





XX DNA involved in streptogramin antibiotic biosynthesis - for  
PT prodn. or bio-conversion of streptogramin(s) or prodn. of  
PT streptogramin intermediates, derivs. or hybrid antibiotics  
XX  
PS Claim 2: Page 54-55; 83pp; French.  
XX  
CC The snac gene product is involved in the biosynthesis of  
CC streptogramins, antibiotics active against Gram-positive bacteria.  
CC The identification of the sequences encoding the enzymes involved  
CC in the biosynthetic pathway means that they can be isolated and  
CC manipulated. Mutant microorganisms in which a step in the  
CC streptogramin biosynthetic pathway is blocked can be cultured to  
CC produce streptogramin intermediates, which may later be converted  
CC to streptogramin derivatives. Recombinant cells may also be used  
CC for the bioconversion of streptogramins from one form to another or  
CC for the production of hybrid antibiotics.  
XX  
SO Sequence 1208 BP; 190 A; 500 C; 360 G; 158 T; 0 other;  
Query Match 49.2%; Score 37.4; DB 15; Length 1208;  
Best Local Similarity 74.6%; Pred. No. 0.017;  
Matches 47; Conservative 0; Mismatches 16; Indels 0; Gaps 0;  
OY 13 ACCGCCGATACGCCACTTTGGCCGTGACAGCGCCGACTTACCTGCGAGGTGTCAG 72  
DB 1114 ACCGCCGCTACGCGCCACTTGGCCGGAATGCGGACTTACCTGCGAGGACCGAC 1173  
OY 73 CCC 75  
DB 1174 CGC 1176  
RESULT 25  
AA064201  
ID AA064201 standard; cDNA: 5392 BP.  
XX  
AC AA064201;  
XX  
DT 18-NOV-1994 (first entry)  
XX  
DE Sequence comprising the snab, snab and snac gene cluster.  
XX  
KM Antibiotic; streptogramin; snab; snab; snac; biosynthesis; enzyme;  
KM biosynthetic pathway; Streptomyces pristinaespiralis; ds.  
XX  
OS Streptomyces pristinaespiralis.  
XX  
PN FR2696189-A.  
XX  
PD 01-APR-1994.  
XX  
PF 25-SEP-1992; 92FR-0011441.  
XX  
PR 25-SEP-1992; 92FR-0011441.  
XX  
PA (RHON ) RHONE POULENC RORER SA.  
XX  
PI Blanc V, Blanche F, Crouzet J, Jacques N, Lacroix P;  
PI Thibaut D, Zagorec M;  
XX  
DR WPI; 1994-128286/16.  
XX  
XX DNA involved in streptogramin antibiotic biosynthesis - for  
PT prodn. or bio-conversion of streptogramin(s) or prodn. of  
PT streptogramin intermediates, derivs. or hybrid antibiotics  
XX  
PS Disclosure; Page 44-47; 83pp; French.  
XX  
CC This sequence comprises the snab, snab and snac genes which are  
CC involved in the biosynthesis of streptogramins, antibiotics active  
CC against Gram-positive bacteria. The identification of the sequences  
CC encoding the enzymes involved in the biosynthetic pathway means that

CC they can be isolated and manipulated. Mutant microorganisms in  
CC which a step in the streptogramin biosynthetic pathway is blocked  
CC can be cultured to produce streptogramin intermediates, which may  
CC later be converted to streptogramin derivatives. Recombinant cells  
CC may also be used for the bioconversion of streptogramins from one  
CC form to another or for the production of hybrid antibiotics.  
XX  
SO Sequence 5392 BP; 811 A; 2161 C; 1671 G; 749 T; 0 other;  
Query Match 49.2%; Score 37.4; DB 15; Length 5392;  
Best Local Similarity 74.6%; Pred. No. 0.02;  
Matches 47; Conservative 0; Mismatches 16; Indels 0; Gaps 0;  
OY 13 ACCGCCGATACGCCACTTTGGCCGTGACAGCGCCGACTTACCTGCGAGGTGTCAG 72  
DB 4671 ACCGCCGCTACGCGCCACTTGGCCGGAATGCGGACTTACCTGCGAGGACCGAC 4730  
OY 73 CCC 75  
DB 4731 CGC 4733  
RESULT 26  
AA199683  
ID AA199683 standard; DNA: 4403765 BP.  
XX  
AC AA199683;  
XX  
DT 15-JAN-2002 (first entry)  
XX  
DE Mycobacterium tuberculosis strain H37Rv genome SEQ ID NO 2.  
XX  
KM Mycobacterium tuberculosis; strain H37Rv; strain CDC 1551; genome;  
KM variation; epidemiology; patient treatment; epidemic monitoring; ds.  
XX  
OS Mycobacterium tuberculosis.  
XX  
PN US6294328-B1.  
XX  
PD 25-SEP-2001.  
XX  
PF 24-JUN-1998; 98US-0103840.  
XX  
PR 24-JUN-1998; 98US-0103840.  
XX  
PA (GENO-) INST GENOMIC RES.  
XX  
PI Fleischmann RD, White OR, Fraser CM, Venter JC;  
XX  
DR WPI; 2001-647261/74.  
XX  
XX Evaluating strain variation of Mycobacterium tuberculosis, comprises  
PT determining the nucleotide sequence of the strain at positions in the  
PT genome corresponding to positions where M. tuberculosis strains CDC  
PT 1551 and H37Rv differ.  
XX  
PS Claim 4: SEQ ID NO 2: 3pp + Sequence Listing; English.  
XX  
XX The invention relates to evaluating strain variation within and between  
CC different populations of the tuberculosis bacterial pathogen,  
CC Mycobacterium tuberculosis or related Mycobacterium by determining the  
CC nucleotide sequence of the first strain at positions in the complete  
CC sequence of the genome that correspond to positions that differ in the  
CC nucleotide sequences of M. tuberculosis strains CDC 1551 (AA199683) and  
CC H37Rv (AA199682). The method is useful for evaluating strain variation of  
CC M. tuberculosis and has valuable application in the fields of  
CC tuberculosis genetics, epidemiology, patient treatment and epidemic  
CC monitoring.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from USPTO  
CC at seqdata.uspto.gov/sequence.html?docID=6294328B1.  
XX  
SO Sequence 4403765 BP; 757105 A; 1447799 C; 1441301 G; 757371 T; 189 other;

Query Match	45.38;	Score 34.4;	DB 22;	Length 4403765;
Best Local Similarity	73.38;	Pred. NO. 0.31;		
Matches 44;	Conservative 0;	Mismatches 16;	Indels 0;	Gaps 0;

QY 12 GAGCGCCGATACGGCCACTTTGGCCGTGACGACGCCGACTTCACCCTCAGAGTGTCGA 71  
| | | | | | | | | | | | | | | | | |  
Db 1567783 GACCGCGCCCTACGGGCACCTTCTGGCCGACCCGACGCTGCAATTATCCCGGAGCACGCTCGA 1567842

RESULT 27  
AAI99682  
ID AAI99682 standard; DNA; 4411529 BP.

AC AAI99682;

DT . 15-JAN-2002 (first entry)

DE Mycobacterium tuberculosis strain H37Rv genome SEQ ID NO 1

KW Mycobacterium tuberculosis; strain H37Rv; strain CDC 1551; genome; variation; epidemiology; patient treatment; epidemic monitoring; ds

OS Mycobacterium tuberculosis.

PN US6294328-B1.

PD 25-SEP-2001.

PF 24-JUN-1998; 98US-0103840.

PR 24-JUN-1998; 98US-0103840.

PA. (GENO-) INST GENOMIC RES.

PI Fleischmann RD, White OR, Fraser CM, Venter JC;

DR WPI; 2001-647261/74.

PT Evaluating strain variation of *Mycobacterium tuberculosis*, comprises  
PT determining the nucleotide sequence of the strain at positions in the  
PT genome corresponding to positions where *M. tuberculosis* strains CDC  
PT 1551 and H37Rv differ -

PS Claim 3; SEQ ID NO 1; 3pp + Sequence Listing; English.

CC The invention relates to evaluating strain variation within and between  
CC different populations of the tuberculosis bacterial pathogen,  
CC *Mycobacterium tuberculosis* or related *Mycobacterium* by determining the  
CC nucleotide sequence of the first strain at positions in the complete  
CC sequence of the genome that correspond to positions that differ in the  
CC nucleotide sequences of *M. tuberculosis* strains CPC 1551 (AM199683) and  
CC H37Rv (AM199682). The method is useful for evaluating strain variation of  
CC *M. tuberculosis* and has valuable application in the fields of  
CC tuberculosis genetics, epidemiology, patient treatment and epidemic  
CC monitoring.

CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from USPTO  
CC at [seqdata.uspto.gov/sequence.html?DocID=6294328B1](http://seqdata.uspto.gov/sequence.html?DocID=6294328B1).

Sequence 4411529 BP; 758565 A; 1449983 C; 1444602 G; 758379 T; 0 other;

Query Match	45.3%	Score 34.4	DB 22	Length 4411529
Best Local Similarity	73.3%	Pred. No. 0.31		
Matches 44	Conservative 0	Mismatches 16	Indels 0	Gaps 0

QY 12 GACGGCCGATTCAGCGCACTTTGGCCGTGACGACGCCGACTTACCCTCCGAGSTGTGTCAA 71  
|||||  
Db 1567944 GACCGCCGCTTAAGGGGACACTTTCGGCCGCACCGACGTCGTGAATTATCCGTGGGACACGCTCGA 1568003

RESULT 28  
AAA81476

ID AAA81476 standard; DNA; 56485 BP.

AC AAAA81476

DT 04-DEC-2000 (first entry)

DE	N. meningitidis partial DNA sequence gnm_24	SEQ ID NO: 24
...	...	...

KW *Neisseria meningitidis*; *Neisseria gonorrhoeae*; genome; immunogenic;  
 KW antigen; vaccine; diagnosis; infection; antibacterial; identification;  
 KW *Meningococcus B*; MenB; ds.

OS *Neisseria meningitidis*

PN WO200022430-A2

PD 20-APR-2000.

PF 08-OCT-1999; 99WO-US23573

PR	09-OCT-1998;	98US-0103794
PR	30-APR-1999;	99US-0132068

PA (CHIR ) GHIRON CORP

PI    Frazer CM,    Hickey E,    Peterson J,    Tettelin H,    Venter JC

PI Rappuoli R, Pizza M;

DR WPI; 2000-318079/27.

PT Isolated nucleotide

PT other Neisserial infections, for example, *N.gonorrhoea* -  
XX  
PS Claim 7; Page 507-524; 1760pp; English.

CC The present invention describes methods

CC represents specifically claimed *Neisseria meningitidis* genomic DNA  
CC sequences: AAA81260 to AAA81303 and AAB25620 to AAB25663 represent  
CC *Neisseria* DNA sequences and their corresponding proteins; AA81254 to  
CC AA81259 and AA81304 to AA81331 represent PCR primers used in the  
CC isolation of *Neisseria meningitidis* DNA sequences; and AA81322 to  
CC AA81452 represent *Neisseria meningitidis* MemB polynucleotide OMP  
CC sequences, which are all used in the exemplification of the present  
CC invention. The nucleic acid sequences, protein sequences, and antipodics  
CC against them, can be used in the manufacture of a composition. The  
CC composition can be used as a medicament (or in the manufacture of a  
CC medicament) for treating, preventing or diagnosing infection due to  
CC *Neisserial* bacteria. For example, some of the identified proteins could  
CC be components of vaccines against *Meningococcus* B; against all serotypes  
CC and/or against all pathogenic *Neisseriae*. Identification of sequences  
CC from the bacterium will also facilitate production of biological probes,  
CC particularly organism-specific probes. Attempts to make efficacious  
CC *Meningococcus* B vaccines have failed mainly due to antigen tolerance.  
CC Multivalent vaccines have also been tried but none have successfully  
CC overcome antigenic variability. The provision of further, complete  
CC sequences may provide an opportunity to identify secreted or surface  
CC exposed proteins that may be presumed targets for the immune system and  
CC which are not antigenically variable or at least more conserved than  
CC other more variable regions.

SQ Sequence 56485 BP; 12504 A; 14247 C; 16158 G; 13573 T; 3 other;

Query Match	43.2%	Score 32.8	DB 21	Length 56485
Best Local Similarity	71.7%	Pred. No. 0.67		
Matches 43: Conservative	0	Mismatches 17	Indels 0	Gaps 0

Oy 4 TTGATCAAGACCGCGGCATACGGCCACTTTTGGCGGTGACGACGCCGACTTCACCCTGGAG 63  
| | | | | | | | | | | | | | | | | |  
Db 54065 TACAGTAATCCGCGCCTTACGGACATTTCGGCGCGAAGAACCTGAGTTTCATTGGGAG 54120

```
RESULT 29
AAAF21612/C
ID   AAAF21612 standard; DNA; 349980 BP.
XX
AC   AAAF21612;
XX
DT   13-MAR-2001 (first entry)
XX
DE   Neisseria meningitidis B nucleotide sequence SEQ ID NO:113.
XX
KW   Neisseria meningitidis; Neisseria gonorrhoeae; immunogenic; vaccine;
KM   diagnosis; antigen; detection; infection; gene therapy; antibacterial;
XX   ds.
XX
OS   Neisseria meningitidis.
XX
PN   WO20006791-A1.
XX
PD   09-NOV-2000.
XX
PF   08-MAR-2000; 2000MO-US05928.
XX
PR   30-APR-1999; 99US-0132068.
PR   08-OCT-1999; 99MO-US23573.
PR   28-FEB-2000; 2000GB-0004695.
XX
PA   (CHIR ) CHIRON CORP.
XX   (GENO-) INST GENOMIC RES.
XX
PI   Pizzo M, Hickey E, Peterson J, Tettelin H, Venter JC, Maignani V,
PI   Galeotti C, Mora M, Ratti G, Scarselli M, Scariato V, Rappuoli R,
PI   Frazer CM, Grandi G;
XX
DR   WPI; 2000-647603/62.
XX
PT   Neisseria meningitidis B full length genome sequence and open reading
PT   frames are used to detect, treat and prevent Neisserial infections -
XX
PS   Claim 7; Appendix A; 692pp; English.
XX
CC   The present invention describes the full length genome of
CC   Neisseria meningitidis B (NMB). The sequences in AAAF21544 and AAAF21607
CC   to AAAF21613 represent fragments of the NMB genomic sequence, as the
CC   sequence was too long to go in a record on its own it was split into 8
CC   sequences which overlap each other at the beginning and end of each
CC   sequence by 49980 bp (i.e. the last 49980 bp of AAAF21544 is repeated at
CC   the beginning of AAAF21607, the last 49980 bp of AAAF21607 are repeated at
CC   the beginning of AAAF21608, and so on). AAAF21545 to AAAF21588 encode the
CC   Neisseria proteins given in AAB58550 to AAB58593, and AAAF21589 to
CC   AAAF21606 represent PCR primers which are used in the exemplification of
CC   the present invention. The NMB genome and fragments from it have
CC   antibacterial activity, and can be used in vaccines and gene therapy.
CC   Neisseria nucleic acids, proteins and/or antibodies which binds to the
CC   proteins can be used in compositions for treating or preventing infection
CC   due to Neisserial bacteria or as a diagnostic reagent for detecting the
CC   presence of Neisserial bacteria or of antibodies raised to Neisserial
CC   bacteria. Computers, computer memory, computer storage medium or computer
CC   databases can be used in a search to identify open reading frames (ORFs)
CC   or coding sequences within the NMB genome. The DNA sequences provide
CC   further opportunities to find antigenic or immunogenic proteins which are
CC   more effective in vaccines than the outer membrane proteins currently
CC   used.
XX
SQ   Sequence 349980 BP; 86473 A; 95646 C; 85908 G; 81953 T; 0 other;
XX
XX
Query Match          43.2%; Score 32.8; DB 21: Length 349980;
Best Local Similarity 71.7%; Pred. No. 0.79;
Matches 43; Conservative 0; Mismatches 17; Indels 0; Gaps 0;
```

```
0Y   4 TTTATCAAGACCGCGCATCTTGCGCGACGACGCGGACGACCTTCGCGAG 63
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db  87588 TACAGTAATCCGCCGCTTACGACATTTTGGCCGCGAAGAACGTGACTTGCGAG 87550
```

```
RESULT 30
AAA81489/C
ID   AAA81489 standard; DNA; 837096 BP.
XX
AC   AAA81489;
XX
DT   04-DEC-2000 (first entry)
XX
DE   N. meningitidis partial DNA sequence gnm_37 SEQ ID NO:37.
XX
KW   Neisseria meningitidis; Neisseria gonorrhoeae; genome; immunogenic;
KM   antigen; vaccine; diagnosis; infection; antibacterial; identification;
XX   Meningococcus B; MenB; ds.
XX
OS   Neisseria meningitidis.
XX
PN   WO200022430-A2.
XX
PD   20-APR-2000.
XX
PF   08-OCT-1999; 99MO-US23573.
XX
PR   09-OCT-1998; 98US-0103794.
PR   30-APR-1999; 99US-0132068.
XX
PA   (CHIR ) CHIRON CORP.
XX
PI   Frazer CM, Hickey E, Peterson J, Tettelin H, Venter JC,
PI   Maignani V, Galeotti C, Mora M, Ratti G, Scarselli M, Scariato V,
PI   Rappuoli R, Pizzo M;
XX
DR   WPI; 2000-318079/27.
XX
PT   Isolated nucleotide sequences of Neisseria meningitidis which can be
PT   used in the diagnosis and treatment of N. meningitidis infection and
PT   other Neisserial infections, for example, N.gonorrhoea.
XX
PS   Claim 7; Page 629-865; 1760pp; English.
XX
CC   The present invention describes methods of obtaining immunogenic
CC   proteins from Neisseria genomic sequences. AAA81453 to AAA82414
CC   represent specifically claimed Neisseria meningitidis genomic DNA
CC   sequences; AAA81260 to AAA81303 and AAB25620 to AAB25663 represent
CC   Neisseria DNA sequences and their corresponding proteins; AAA81254 to
CC   AAA81259 and AAA81304 to AAA81321 represent PCR primers used in the
CC   isolation of Neisseria meningitidis DNA sequences; and AAA81322 to
CC   AAA81452 represent Neisseria meningitidis MenB polynucleotide ORF
CC   sequences, which are all used in the exemplification of the present
CC   invention. The nucleic acid sequences, protein sequences, and antibodies
CC   against them, can be used in the manufacture of a composition. The
CC   composition can be used as a medicament (or in the manufacture of a
CC   medicament) for treating, preventing or diagnosing infection due to
CC   Neisserial bacteria. For example, some of the identified proteins could
CC   be components of vaccines against Meningococcus B; against all serotypes;
CC   and/or against all pathogenic Neisseriae. Identification of sequences
CC   from the bacterium will also facilitate production of biological probes,
CC   particularly organism-specific probes. Attempts to make efficacious
CC   Meningococcus B vaccines have failed mainly due to antigen tolerance.
CC   Multivalent vaccines have also been tried but none have successfully
CC   overcome antigenic variability. The provision of further, complete
CC   sequences may provide an opportunity to identify secreted or surface
CC   exposed proteins that may be presumed targets for the immune system and
CC   which are not antigenically variable or at least more conserved than
CC   other more variable regions.
XX
SQ   Sequence 837096 BP; 207534 A; 227065 C; 205215 G; 197280 T; 2 other;
XX
XX
Query Match          43.2%; Score 32.8; DB 21: Length 837096;
Best Local Similarity 71.7%; Pred. No. 0.85;
Matches 43; Conservative 0; Mismatches 17; Indels 0; Gaps 0;
```



S0 Sequence 657 BP; 155:A; 175 C; 134 G; 193 T; 0 other;

Query Match 37.6%; Score 28.6; DB 21; Length 657;  
Best Local Similarity 61.3%; Pred. No. 9;  
Matches 46; Conservative 0; Mismatches 29; Indels 0; Gaps 0;

OY 1 AGGTCATCAACACGGCCCATACGGCCACTTTGGGCCCGTAGCAGACGCCGCACTTCACTTCG 60  
|| ||| | ||||| | ||| |||| | |||| ||| ||| | ||| ||  
Db 221 ATGGGCATTGACGACGCCGAATAATGTGGCGATTGGCTTAATGACTACGTGCAATTACC CGC 280

OY 61 GAGGTGTCGAAGCCC 75  
|| ||| ||| ||  
Db 281 GACGTGATCAGTGCC 295

RESULT 33

AAF08609  
ID AAF08609 standard; cDNA; 383 BP.

XX AAF08609;  
XX  
XX 13-MAR-2001 (first entry)  
DE Fusarium venenatum EST SEQ ID NO:1132.  
XX  
XX Fusarium venenatum EST SEQ ID NO:1132.  
KW Multiple gene expression; filamentous fungal cell; EST;  
KM expressed sequence tag; Fusarium venenatum; Aspergillus niger;  
RV Aspergillus oryzae; Trichoderma reesei; identification; recombinaton;  
KW culture condition; environmental stress; spore morphogenesis;  
KM metabolic pathway engineering; catabolic pathway engineering; ss.  
XX  
OS Fusarium venenatum.  
XX  
XX WO200056762-A2.  
PN  
PD 28-SEP-2000.  
XX  
PF 22-MAR-2000; 2000WO-US07781.  
PR  
PR 22-MAR-1999; 99US-0273623.  
XX  
XX (NOVO ) NOVO NORDISK BIOTECH INC.  
PA (NOVO ) NOVO NORDISK AS.  
XX  
PI Berka RM, Rey MW, Shuster JR, Kauppinen S, Clausen IG, Olsen PB;  
XX WPI: 2000-594572/56.  
DR  
XX  
PT Monitoring differential expression of genes in filamentous fungal cells  
PT uses fluorescence-labeled nucleic acids isolated from the cells and a  
XX substrate of expressed sequence tags -  
PS  
PS Claim 86; Page 813; 3161pp; English.  
XX

The present invention describes a method for monitoring differential expression of genes in a first filamentous fungal (FF) cell relative to expression of the same genes in one or more second filamentous fungal cells. The method uses fluorescence-labeled nucleic acids isolated from the FF cells and a substrate of expressed sequence tags (EST). The ESTs are used in the methods for monitoring differential expression of genes in a first filamentous fungal (FF) cell relative to expression of the same genes in one or more second filamentous fungal cells. Monitoring potential of the microorganisms to be improved. New genes may be discovered, possible functions of unknown open reading frames can be identified and gene copy number variation and stability can be monitored. The expression of genes can be used to study how FF cells adapt to changes in culture conditions, environmental stress, spore morphogenesis, recombinaton, metabolic or catabolic pathway engineering. Using ESTs provides several advantages over genomic or random cDNA clones including elimination of redundancy as one spot on an array equals one gene or open reading frame, and organisation of the microarrays based on function of the gene products to facilitate

CC	analysis of the results. AAF07478 to AAP11247 represents ESTs from
CC	Fusarium venenatum; AAFL1248 to AAP11853 represents ESTs from Aspergillus
CC	niger; AAP11854 to AAFL4878 represents ESTs from Aspergillus oryzae; and
CC	AAFL4879 to AAP15337 represents ESTs from Trichoderma reesei, which are
CC	all specifically claimed in the present invention.
XQ	Sequence 383 BP; 74 A; 115 C; 87 G; 89 T; 18 other;
	Query Match 36.6%; Score 27.8; DB 21; Length 383;
	Best Local Similarity 61.1%; Pred. No. 15;
	Matches 44; Conservative 0; Mismatches 28; Indels 0; Gaps 0;
OY	4 TTCATCAAGACCGGCGCATCGGCACCTTTGGCCGTGACACGCCGCACTTACTGCGAG 63
Db	145 TACATTTGAGAACCCTGCTCCTGTCNACTTTGCCCGCGCGGCGCTGCGCACCGAGAC 204
OY	64 GTGGTCAAGCCC 75
Db	205 TTCTTTCATTTCCC 216
RESULT 34	
ABK13571	
ID	ABK13571 standard; cDNA; 2284 BP.
XX	ABK13571:
XX	23-APR-2002 (first entry)
XX	Ryegrass 4-coumarate Co-A-ligase 1 (Lp4CL1) cDNA.
DE	
KW	Perennial ryegrass; ss; lignin; 4 coumarate COA-ligase; QTL; gene;
KM	lignin biosynthesis; enzyme; cinamoyl-CoA reductase; CCR; Lp4CL1;
KM	cinamoyl alcohol dehydrogenase; CAD; molecular genetic marker;
KM	qualitative trait loci; tagging; QTL mapping; DNA fingerprinting;
KM	marker assisted selection; forage improvement; turf grass improvement;
KW	dry matter digestibility; herbage quality; palatability; regrowth;
KW	cold tolerance; drought tolerance; tiller survival; plant persistence.
XX	
OS	Lolium perenne.
XX	
Key	Location/Qualifiers
FT	1..322
FT	/tag= a
FT	323..2146
FT	/tag= b
FT	/product= "LP4CL1 protein"
FT	/transl_except= (pos:2030..2038, aa:Ala Arg)
FT	3'UTR 2147..2266
FT	/tag= c
FT	2267..2284
FT	/tag= d
XX	
PN	MO200195702-Al.
XX	
PD	20-DEC-2001.
PF	14-JUN-2001; 2001WO-AU00699.
PR	14-JUN-2000; 2000AU-0008154.
XX	
PA	(VIC-) STATE VICTORIA DEPT NATURAL RES & ENVIRO.
PA	(UVAD-) UNIV ADELAIDE.
PA	(ITMA-) INT MAIZE & WHEAT IMPROVEMENT CENT.
PA	(SAUS-) STATE SOUTH AUSTRALIA SOUTH AUSTRALIAN R.
PA	(USC-) UNIV SOUTHERN CROSS.
PA	(DAIR-) DAIRY RES & DEV CORP.
XX	
PI	Spengenberg GC, Lidgett AJ, Heath RL, McInnes RL, Lynch DP;
XX	
DR	WPI: 2002-097993/13.
XR	P-FSDB: AAU75084.
XX	

PT Novel nucleic acid encoding enzymes involved in lignin biosynthetic  
 PT pathway from ryegrass or fescue species useful for modifying lignin  
 PT biosynthesis in plants and as a molecular genetic marker  
 XX  
 PS Claim 3; Fig 2; 148bp; English.

CC This invention represents purified or isolated nucleic acid and protein  
 CC sequences of enzymes involved in lignin biosynthesis. The enzymes  
 CC of the invention are 4 coumarate CoA-ligase (4CL), cinnamoyl-CoA  
 CC reductase (CCR) and cinnamyl alcohol dehydrogenase (CAD) from a ryegrass  
 CC (*Lolium sp.*) or fescue (*Festuca sp.*). The invention also comprises an  
 CC isolated regulatory element from the nucleic acid sequences and a plant  
 CC cell or seed transformed with the nucleic acid. An isolated regulatory  
 CC element from these nucleotide molecules is useful for expressing an  
 CC exogenous gene in plant cells. The nucleotide sequences of the invention  
 CC and vectors containing these sequences are useful for modifying lignin  
 CC biosynthesis in a plant and are useful as a molecular genetic marker for  
 CC qualitative trait loci (QTL) tagging, QTL mapping, DNA fingerprinting,  
 CC and in marker assisted selection, in forage and turf grass improvement,  
 CC e.g. tagging QTLs for dry matter digestibility, herbage quality,  
 CC palatability, regrowth after cutting and grazing, cold tolerance,  
 CC drought tolerance, tiller survival and plant persistence. The present  
 CC sequence represents the perennial ryegrass 4-coumarate Co-A-ligase 1  
 CC (Lp4CL1) cDNA of the invention.

XX  
 SQ Sequence 2284 BP; 391 A; 750 C; 733 G; 410 T; 0 other;

Query Match 36.6%; Score 27.8; DB 24; Length 2284;  
 Best Local Similarity 62.0%; Pred. No. 18;  
 Matches 44; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

OY 2 GGTTCATCAGACCGCGCATACGCGCCTTTGGCCGTGACGACCGGACTTTCACCTGCG 61  
 DB 1647 GGTGGCTCCACACGCGGACATCGCTACGTCGACGACGACGAGGCTTTCATCGCG 1706  
 OY 62 AGGTGTCACG 72  
 DB 1707 ACCGCGTCACG 1717

RESULT 35  
 AAT13227  
 ID AAT13227 standard; DNA; 3600 BP.  
 AC AAT13227;  
 DT 16-SEP-1996 (first entry)  
 DE Thermostable enzyme (converting maltose to trehalose) DNA.  
 XX  
 KW Thermostable enzyme; thermophilic bacterium; sweetener; trehalose;  
 KM maltose; food; recombinant enzyme; cosmetic; pharmaceutical; ds.  
 XX  
 OS *Thermus aquaticus* ATCC 33923.  
 XX  
 FH Key Location/Qualifiers  
 FT 5'UTR 1..540  
 FT /\*tag= a  
 FT mat\_peptide 541..3429  
 FT /\*tag= b  
 FT 3'UTR 3430..3600  
 FT /\*tag= c  
 XX  
 PN EP704531-A2.  
 PD 03-APR-1996.  
 XX  
 PF 29-SEP-1995; 95EP-0306875.  
 XX  
 PR 08-SEP-1995; 95JP-0255829.  
 PR 01-OCT-1994; 94JP-0260984.  
 XX  
 PA (HAYB ) HAYASHIBARA SEIBUTSU KAGAKU.

XX  
 PI Kubota M, Sugimoto T, Tsusaki K;  
 XX  
 DR WPI: 1996-173035/18.  
 DR P-PSDB: AAW9181, AAW9182.  
 XX  
 PT Isolated DNA encoding enzyme for converting maltose to trehalose  
 PT used for prodn. of trehalose for use in food prods., cosmetics and  
 PT pharmaceuticals, partic. as sweetener  
 XX  
 PS Claim 8; Page 28-31; 46pp; English.

CC This DNA encoding a thermostable enzyme which converts maltose to  
 CC trehalose may be used to express the enzyme recombinantly in *E. coli*  
 CC using plasmid vector Bluescript II SK(+) or plasmid pKK223-3. The  
 CC enzyme is then produced by culturing the transformant. The enzyme  
 CC catalyses a reaction to produce high yields of trehalose at high  
 CC temp., which prevents bacterial contamination. Trehalose is used in  
 CC food products, cosmetics and pharmaceuticals, as a sweetener, etc.,  
 XX

XX  
 SQ Sequence 3600 BP; 559 A; 1346 C; 1168 G; 527 T; 0 other;

Query Match 36.6%; Score 27.8; DB 17; Length 3600;  
 Best Local Similarity 62.0%; Pred. No. 19;  
 Matches 44; Conservative 0; Mismatches 27; Indels 0; Gaps 0;

OY 2 GGTTCATCAGACCGCGCATACGCGCCTTTGGCCGTGACGACCGGACTTTCACCTGCG 61  
 DB 1412 GGATCCCGGAACACCGCCAGTGGCCCTCTTCCTCGACACGACGAGCTCACCTCGG 1471  
 OY 62 AGGTGTCACG 72  
 DB 1472 AGAAGTCACG 1482

RESULT 36  
 AAX19362  
 ID AAX19362 standard; DNA; 7584 BP.  
 AC AAX19362;  
 DT 19-MAY-1999 (first entry)  
 DE Rhodococcus corallina ohp operon.  
 XX  
 KW Rhodococcus corallina: ohp operon; biosensor; mycolic acid bacteria;  
 KM inducible promoter; environmental pollutant; industry; medicine; ds.  
 XX  
 OS Rhodococcus corallina.  
 XX  
 PN WO9900517-A2.  
 PD 07-JAN-1999.  
 XX  
 PF 29-JUN-1998; 98WO-GB01893.  
 XX  
 PR 27-JUN-1997; 97GB-0013666.  
 XX  
 PA (UYCA-) UNIV CAMBRIDGE TECH SERVICES LTD.  
 XX  
 PI Archer JAC, Powell JAC, Roland HU, Summers DK;  
 XX  
 DR WPI: 1999-095760/08.  
 DR P-PSDB: AAW9181, AAW9182, AAW9183, AAW9184, AAW9185, AAW9186.  
 XX  
 PT Isolating DNA encoding inducible promoter from mycolic acid bacteria  
 PT - useful to produce mycolic acid bacterial biosensors for particular  
 PT analyses, such as environmental pollutants, e.g. from industry or  
 PT medicine  
 XX  
 PS Example 7; Fig 4; 67pp; English.  
 CC A method has been developed for identifying and/or isolating DNA from





```

XX 15-JUL-1998; 98US-0092866.
XX (DUPO ) DU PONT DE NEMOURS & CO E I.
XX (PION-) PIONEER HI-BRED INT INC.
XX Famodu LO, Orozco EM, Rafalski JA;
XX WPI: 2001-3388927/41.
XX P-PSDB: AAE03590.
XX New isolated polynucleotide encoding an aspartyl-L-lysine synthetase useful
XX as targets to facilitate design and/or identification of inhibitors of
XX those enzymes that may be useful as herbicides -
XX Example 4; Column 39-42; 40pp; English.
XX The present sequence is a cDNA encoding rice cysteinyl-L-lysine synthetase
XX of the invention. The cysteinyl-L-lysine synthetase are used as targets to
XX facilitate designing and identification of inhibitors of the enzymes
XX which are useful as herbicides. All or a substantial portion of the
XX nucleic acid fragments of the present invention are used as probes for
XX genetically and physically mapping the genes that they are a part of,
XX and as markers for traits linked to those genes. Such information is
XX useful in plant breeding in order to develop lines with desired
XX phenotypes.
XX Sequence 1957 BP; 599 A; 384 C; 470 G; 504 T; 0 other:
XX
XX Query Match 36.3%; Score 27.6; DB 22; Length 1957;
XX Best Local Similarity 63.6%; Pred. No. 20;
XX Matches 42; Conservative 0; Mismatches 24; Indels 0; Gaps 0;
XX
OY 6 CATCAAGACCGCCGATACGGCCTTTGGCGGACGCGGACTGACCTGCGAGT 65
DB 176 GGTACGCGCTTACGCTTACGCGACATGCGCGCGCGCTTACGCTTACGAGT 235
OY 66 GGTCAA 71
DB 236 CCTCTA 241

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XX Novel DNA array useful for determining differential expression of
XX Methylococcus capsulatus genes, comprises polynucleotides or
XX oligonucleotides representative for a selective number of Methylococcus
XX capsulatus genes -
XX Claim 14; Page 296-297; 678pp; English.
XX The invention relates to a novel DNA array giving a representation of a
XX number of Methylococcus capsulatus genes. The method of the invention is
XX useful for determination of the differential expression of the genes of
XX M. capsulatus, and for studying gene expression on a genomic scale and in
XX gene expression assays of M. capsulatus genes. The sequences shown in
XX AB090016-AB091855 represent M. capsulatus genes for use in arrays of the
XX invention.
XX Sequence 2907 BP; 504 A; 975 C; 887 G; 541 T; 0 other:
XX
XX Query Match 36.3%; Score 27.6; DB 24; Length 2907;
XX Best Local Similarity 60.8%; Pred. No. 21;
XX Matches 45; Conservative 0; Mismatches 29; Indels 0; Gaps 0;
XX
OY 3 GTTCATCAAGACCGCCGATACGGCCTTTGGCGGACGCGGACTTACCTGCGA 62
DB 915 GTTCATCAATGATCGCCCAATATGCGACATGCGCGCTTACCTGCGTGA 974
OY 63 GGTGTCAAACCGCC 76
DB 975 CGGCGAAGAACGCC 988

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RESULT 40
AA225727
ID AA225727 standard; cDNA: 1791 BP.
AC AA225727;
XX 05-JAN-2000 (first entry)
DE Stachybotrys chartarum phenol oxidising enzyme encoding cDNA.
XX Stachybotrys chartarum; phenol oxidising enzyme; colour; dye;
XX detergent; anti-dye transfer; stain removal; bleaching; ss.
XX Stachybotrys chartarum.
XX OS
XX WO9949010-A2.
XX PN
XX 30-SEP-1999.
XX 23-MAR-1999; 99WO-EP02042.
XX PF
XX 24-MAR-1998; 98US-0046969.
XX PR
XX 22-DEC-1998; 98US-0218702.
XX (UNIL ) UNILEVER NV.
XX (UNIL ) UNILEVER PLC.
XX
XX Convents D, Amory A, Wang H, Dhaese P, Lambrechts-Rongvaux A;
XX Wang C;
XX WPI: 1999-601211/51.
XX P-PSDB: AAV45222.
XX
XX Detergent composition containing phenol oxidase from Stachybotrys, used
XX to bleach stains and prevent dye transfer -
XX Example 15; Fig 5; 56pp; English.
XX The present invention describes a detergent composition containing a
XX purified phenol oxidising enzyme derived from Stachybotrys. The present
XX sequence encodes Stachybotrys chartarum phenol oxidising enzyme. The
XX enzyme can be used to modify the colour of dyes and other coloured

```

CC compounds (e.g. for use in pulp and paper bleaching also for removing  
CC stains, e.g. food, tea, blood etc., from fabrics) and for preventing dye  
CC transfer during fabric washing.

XX Sequence 1791 BP; 380 A; 554 C; 448 G; 409 T; 0 other;

#### Query Match

Best Local Similarity 36.1%; Score 27.4; DB 20; Length 1791;

Matches 43; Conservative 0; Mismatches 26; Indels 0; Gaps 0;

OY 4 TTCATCAGACGCCGACATCTTGCCGCTGACGACGCGACTTCACTTCGCGAG 63

DB 577 TTCATCAGACGCTGAGATGCTTGTGTCAGGCTGGCCCTACATTATCAACGAC 636

OY 64 GTGCTCAAG 72

DB 637 GAGGCTGAG 645

#### RESULT 41

AAZ27601

ID AAZ27601 standard; DNA; 1791 BP.

XX AAZ27601;

DT 16-DEC-1999 (first entry)

DE Stachybotrys phenol oxidase coding sequence.

KW Phenol oxidase; enzyme; coloured compound; dye transfer prevention;  
KM fabric washing; stain bleaching; anti-dye transfer; detergent; ss.

XX Stachybotrys chartarum.

XX WO9949020-A2.

XX 30-SEP-1999.

XX 23-MAR-1999; 99WO-US06327.

XX 24-MAR-1998; 98US-0046969.

XX 22-DEC-1998; 98US-0218702.

XX 22-MAR-1999; 99US-0273957.

XX (GENEV) GENENCOR INT INC.

XX Amory A, Wang H, Dhase P, Lambrechts-Rongvaux A, Wang C;

XX WPI: 1999-591088/50.

XX P-PSDB: AAY39992.

XX Novel enzyme for modifying coloured compounds used to prevent  
PT dye-transfer.

XX Claim 21; Fig 5; 64pp; English.

CC This sequence encodes the Stachybotrys chartarum phenol oxidase enzyme  
CC of the invention. The invention is used to modify a coloured compound and  
CC prevent dye transfer during fabric washing, or for stain bleaching or  
CC anti-dye transfer. It is useful in the detergent, paper and pulp, textile  
CC and food industries.

XX Sequence 1791 BP; 380 A; 551 C; 451 G; 409 T; 0 other;

#### Query Match

Best Local Similarity 36.1%; Score 27.4; DB 20; Length 1791;

Matches 43; Conservative 0; Mismatches 26; Indels 0; Gaps 0;

OY 4 TTCATCAGACGCCGACATCTTGCCGCTGACGACGCGACTTCACTTCGCGAG 63

DB 577 TTCATCAGACGCTGAGATGCTTGTGTCAGGCTGGCCCTACATTATCAACGAC 636

OY 64 GTGCTCAAG 72

DB 637 GAGGCTGAG 645

#### RESULT 42

AA50019

ID AA50019 standard; DNA; 1791 BP.

XX AA50019;

DT 10-OCT-2000 (first entry)

DE Stachybotrys chartarum phenol oxidising enzyme cDNA.

KW Phenol oxidising enzyme; detergent; bleaching; ss.

XX Stachybotrys chartarum.

XX Key Location/Qualifiers

XX CDS 7..1791

XX /tag= a

XX WO200039306-A2.

XX 06-JUL-2000.

XX 20-DEC-1999; 99WO-EPI0287.

XX 23-DEC-1998; 98US-0220871.

XX 23-JUN-1999; 99US-0338723.

XX (UNIL) UNILEVER NV.

XX (UNIL) UNILEVER PLC.

XX (HIND-) HINDUSTAN LEVER LTD.

XX Bodie EA, Van Der Velden S, De Vries CH, Wang H;

XX WPI: 2000-514528/46.

XX P-PSDB: AAY95537.

XX Detergent composition comprising novel phenol oxidising enzyme obtained  
PT from fungus or bacteria, useful for pulp and paper bleaching, bleaching  
PT color of stains on fabric and for anti-dye redeposition

XX Disclosure; Fig 5A-B; 45pp; English.

CC The present sequence is that of the Stachybotrys chartarum MUCL 38898  
CC cDNA encoding a phenol oxidising enzyme (see AAY95537). The invention  
CC relates to detergent compositions comprising novel phenol oxidising  
CC enzymes that are encoded by nucleic acids capable of hybridising to  
CC the S. chartarum phenol oxidising enzyme gene (see AA50019), provided  
CC the enzymes are capable of modifying the colour associated with dyes  
CC or coloured compounds, and are produced from a bacterium, yeast or  
CC fungus (see AAY9538-40). The phenol oxidising enzymes can be used  
CC for pulp and paper bleaching, for bleaching the colour of stains on  
CC fabric and for anti-dye transfer in detergent and textile  
CC applications. They may also be capable of modifying the colour in  
CC the absence or presence of an enhancer. Expression vectors and host  
CC cells comprising a nucleic acid encoding a phenol oxidising enzyme,  
CC methods for producing the phenol oxidising enzyme, and methods for  
CC constructing expression hosts are provided.

XX Sequence 1791 BP; 380 A; 551 C; 451 G; 409 T; 0 other;

#### Query Match

Best Local Similarity 36.1%; Score 27.4; DB 21; Length 1791;

Matches 43; Conservative 0; Mismatches 26; Indels 0; Gaps 0;

OY 4 TTCATCAGACGCCGACATCTTGCCGCTGACGACGCGACTTCACTTCGCGAG 63

DB 577 TTCATCAGACGCTGAGATGCTTGTGTCAGGCTGGCCCTACATTATCAACGAC 636

OY 64 GTGCTCAAG 72

Db 637 GAGGCTGAG 645

## RESULT 43

AA51314  
ID AAA51314 standard; DNA: 1791 BP.

XX  
AC AAA51314;

DT 09-OCT-2000 (first entry)

XX Stachybotrys chartarum phenol oxidizing enzyme cDNA.

XX Phenol oxidizing enzyme: colour; dye; modification; detergent; stain;

KW pulp; paper bleaching; ss.

XX Stachybotrys chartarum.

XX Key Location/Qualifiers

FT CDS 7..1791

FT /tag= a

FT /product= Phenol\_Oxidizing\_Enzyme

PN MO200037654-A2.

PD 29-JUN-2000.

PE 20-DEC-1999; 99MO-US31009.

XX 23-DEC-1998; 98US-0220871.

PR 23-JUN-1999; 99US-0338723.

XX (GENMV ) GENENCOR INT INC.

PI Wang H, Bodie EA;

XX WPI: 2000-452191/39.

DR P-PSDB: AAY96761.

XX New phenol oxidizing enzyme for modifying colors associated with dyes

PT or colored compounds, is obtained from fungus and is encoded by a

PT nucleic acid comprising a specific nucleotide sequence

XX Disclosure: Fig 5A-B; 45pp; English.

XX This cDNA encodes Stachybotrys chartarum phenol oxidizing enzyme.

CC Phenol oxidizing enzymes encoded by nucleic acid sequences which

CC hybridize to this DNA are claimed, as long as the enzyme is capable of

CC modifying the colour associated with dyes or coloured compounds. The

CC enzymes are useful in detergent compositions and for modifying colors

CC associated with dyes or colored compounds which occur in stains in a

CC sample. The enzymes are also useful for pulp and paper bleaching,

CC anti-dye transfer in detergent and other textile applications.

XX Sequence 1791 BP; 380 A; 551 C; 451 G; 409 T; 0 other:

## RESULT 44

AA47584  
ID AAL47584 standard; cDNA: 1791 BP.

XX  
AC AAL47584;

DT 13-SEP-2002 (first entry)

XX S chartarum phenol oxidizing enzyme cDNA.

XX Phenol oxidizing enzyme: enzyme; fungus; redox reaction; detergent;

XX paper industry; pulp industry; textile; food industry; gene; ss.

XX Stachybotrys chartarum.

XX Key Location/Qualifiers

FT CDS 7..1791

FT /tag= a

FT /product= "phenol oxidizing enzyme"

PN US6399329-B1.

PD 04-JUN-2002.

PE 21-DEC-1999; 99US-0468578.

XX 12-DEC-1998; 98US-0220871.

PR 23-JUN-1999; 99US-0338723.

XX (GENMV ) GENENCOR INT INC.

PI Wang H, Bodie EA;

XX WPI: 2002-498835/53.

DR P-PSDB: AAO18210.

XX New polynucleotides encoding phenol oxidizing enzymes, useful for

PT preventing the transfer of dyes in solution from one textile to another

PT during detergent washing -

XX Disclosure: Fig 5; 37pp; English.

XX The present invention provides the protein and coding sequences of phenol

CC oxidizing enzymes from Stachybotrys chartarum, Bipolaris spicifera and

CC Curvularia pallescens. These enzymes are useful in the textiles, paper,

CC pulp, detergent and food industries. In particular they are useful for

CC preventing the transfer of dyes in solution from one textile to another

CC during detergent washing (dye transfer inhibition). The present sequence

CC is the S. chartarum phenol oxidizing enzyme cDNA.

XX Sequence 1791 BP; 380 A; 551 C; 451 G; 409 T; 0 other:

XX Query Match 36.1%; Score 27.4; DB 24; Length 1791;

XX Best Local Similarity 62.3%; Pred. No. 23;

XX Matches 43; Conservative 0; Mismatches 26; Indels 0; Gaps 0;

OY 4 TTCATCAAGACCGCGCATATGCGCCGCTTGAGAGCGCGCATTCACCTGCGAG 63

DB 577 TTCATCAAGACCGCGCATATGCGCCGCTTGAGAGCGCGCATTCACCTGCGAG 636

OY 64 GTGCTCAAG 72

DB 637 GAGGCTGAG 645

## RESULT 45

ABA92911  
ID ABA92911 standard; DNA: 7248 BP.

XX  
AC ABA92911;

DT 09-APR-2002 (first entry)

XX Stachybotrys chartarum laccase gene.

XX Laccase; enzyme; blue copper oxidase; fungal; oxidation; reduction;

KM copper; metal cofactor; gene; plant; ds.  
 XX  
 OS Stachybotrys chartarum.  
 XX  
 PN WO200196543-A2.  
 XX  
 PD 20-DEC-2001.  
 XX  
 PF 14-JUN-2001; 2001WO-US19174.  
 XX  
 PR 15-JUN-2000; 2000US-211732P.  
 XX  
 PA (PROD-) PRODIGENE INC.  
 PA (GENY) GENENCOR INC.  
 XX  
 PI Hood E, Howard JA, Bailey M, Van Gastel FJC, Ward M, Wang H;  
 PI Woodard S;  
 XX  
 DR WPI; 2002-090204/12.  
 XX  
 PT Improving recovery of active enzyme e.g. laccase, which requires  
 PT transitional metal cofactor e.g. copper for activity, from a plant, by  
 PT introducing plant nucleotide sequences encoding the enzyme and exposing  
 PT it to cofactor  
 XX  
 PS Example 5; Fig 16A-E; 81pp; English.  
 XX  
 CC The present invention describes a method for improving the recovery of  
 CC an active enzyme from a plant where the enzyme requires a transitional  
 CC metal cofactor for activity. The method comprises introducing into the  
 CC plant nucleotide sequences encoding the enzyme and exposing the enzyme  
 CC to the metal cofactor. The method is useful for improving recovery of  
 CC active enzyme which requires a transitional metal cofactor for activity,  
 CC preferably for improving recovery of active laccase which requires  
 CC copper for activity. The method can be used for improving recovery of  
 CC active organophosphate hydrolase (OPH, E.C. 3.1.8.1) which requires  
 CC zinc, nickel, cobalt or manganese for activity, where the method further  
 CC comprises adding bicarbonate ion salt. The present sequence encodes the  
 CC fungal Stachybotrys chartarum laccase enzyme. Laccases are also called  
 CC blue copper oxidases and use copper to accept and donate electrons in  
 CC the oxidation and reduction of substrates.  
 XX  
 SQ Sequence 7248 BP; 1928 A; 1744 C; 1579 G; 1997 T; 0 other;  
 Query Match 36.1%; Score 27.4; DB 24; Length 7248;  
 Best Local Similarity 62.3%; Pred. No. 26;  
 Matches 43; Conservative 0; Mismatches 26; Indels 0; Gaps 0;  
 OY 4 TTCATCAAGACGGCGCATATAGCGGCGACTTGGCGGTGACGACGGCGACTTCACCTGGAG 63  
 DB 5768 TTCATCAAGACGTGCTGAGAAATGCTACTTGTGACGTGGCGGCTTATATCAACGAC 5827  
 OY 64 GTGGTCAAG 72  
 DB 5828 GACGCTGAG 5836

Search completed: April 23, 2003, 15:42:04  
 Job time : 2439.94 secs



Db 382 GluValValIysPro 386  
 RESULT 2  
 P00817  
 methionine adenosyltransferase (EC 2.5.1.6) - rape (fragment)  
 C:Species: Brassica napus (rape)  
 C:Date: 03-May-1994 #sequence\_revision 07-Oct-1994 #text\_change 05-May-2000  
 C:Accession: P00817  
 R:Park, Y.S.; Kwak, J.M.; Kwon, O.Y.; Kim, Y.S.; Lee, D.S.; Cho, M.J.; Lee, H.H.; Nam, H.  
 Plant Physiol. 103, 359-370, 1993  
 A:Title: Generation of expressed sequence tags of random root cDNA clones of Brassica na  
 A:Reference number: P00816; MUID:94302145; PMID:8029332  
 A:Accession: P00817  
 A:Molecule type: mRNA  
 A:Residues: 1-68 <PAR>  
 A:Experimental source: root, cv. Naehan  
 C:Superfamily: methionine adenosyltransferase  
 C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
 Pred. No.: 1.31e-09 Length: 68  
 Score: 120.00 Matches: 22  
 Percent Similarity: 92.00% Conservative: 1  
 Best Local Similarity: 88.00% Mismatches: 2  
 Query Match: 86.96% Indels: 0  
 DB: 2 Gaps: 0

US-09-198-779B-1\_COPY\_160\_235 (1-76) x P00817 (1-68)

OY 1 AGTTCATCAAGACCGCCGATACGGCCACTTGGCCGTCAGACGCCGACCTTCACCTGC 60  
 Db 36 ArgPheLeuLysThrAlaIatyrGlyHisPheGlyArgAspAspProAspPheThrTrp 55

OY 61 GAGGTGTCACAGCC 75  
 Db 56 GluValValIysPro 60

RESULT 3  
 JN0131  
 methionine adenosyltransferase (EC 2.5.1.6) - Arabidopsis thaliana  
 N:Alternate names: S-adenosylmethionine synthetase  
 C:Species: Arabidopsis thaliana (mouse-ear cress)  
 C:Date: 30-Sep-1991 #sequence\_revision 30-Sep-1991 #text\_change 05-May-2000  
 C:Accession: JN0131  
 R:Peleman, J.; Boerjan, W.; Engler, G.; Seurinck, J.; Bolterman, J.; Alliotte, T.; Van M  
 Plant Cell 1, 81-93, 1989  
 A:Title: Strong cellular preference in the expression of a housekeeping gene of Arabidop  
 A:Reference number: JN0131; MUID:92386056; PMID:2535470  
 A:Accession: JN0131  
 A:Molecule type: DNA  
 A:Residues: 1-393 <PEL>  
 A:Cross-references: GB:M55077; NID:q166871; PIDN:AAA32868.1; PID:q166872  
 A:Experimental source: var. R85  
 A:Note: the sequence derived from var. Columbia differs from that shown in having 117-Gl  
 A:Note: the authors translated the codon GAC for residue 117 as Glu  
 C:Comment: S-Adenosylmethionine synthetase catalyzes the biosynthesis of adenosylmethio  
 C:Genetics:  
 A:Gene: sam-1  
 C:Superfamily: methionine adenosyltransferase  
 C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
 Pred. No.: 1.31e-09 Length: 393  
 Score: 120.00 Matches: 22  
 Percent Similarity: 92.00% Conservative: 1  
 Best Local Similarity: 88.00% Mismatches: 2  
 Query Match: 86.96% Indels: 0  
 DB: 2 Gaps: 0

US-09-198-779B-1\_COPY\_160\_235 (1-76) x JN0131 (1-393)

OY 1 AGTTCATCAAGACCGCCGATACGGCCACTTGGCCGTCAGACGCCGACCTTCACCTGC 60

Db 361 ArgPheLeuLysThrAlaIatyrGlyHisPheGlyArgAspAspProAspPheThrTrp 380  
 OY 61 GAGGTGTCACAGCC 75  
 Db 381 GluValValIysPro 385

RESULT 4  
 S38875  
 methionine adenosyltransferase (EC 2.5.1.6) - tomato  
 N:Alternate names: S-adenosyl-L-methionine synthetase  
 C:Species: Lycopersicon esculentum (tomato)  
 C:Date: 22-Jan-1994 #sequence\_revision 10-Nov-1995 #text\_change 05-May-2000  
 C:Accession: S46539; S38875  
 R:Espartero, J.; Pintor-Toro, J.A.; Pardo, J.M.  
 Plant Mol. Biol. 25, 217-227, 1994  
 A:Title: Differential accumulation of S-adenosylmethionine synthetase transcripts in  
 A:Reference number: S46538; MUID:94289646; PMID:8018871  
 A:Accession: S46539  
 A:Status: nucleic acid sequence not shown  
 A:Molecule type: mRNA  
 A:Residues: 1-393 <ES2>

Alignment Scores:  
 Pred. No.: 1.31e-09 Length: 393  
 Score: 120.00 Matches: 22  
 Percent Similarity: 92.00% Conservative: 1  
 Best Local Similarity: 88.00% Mismatches: 2  
 Query Match: 86.96% Indels: 0  
 DB: 2 Gaps: 0

US-09-198-779B-1\_COPY\_160\_235 (1-76) x S38875 (1-393)

OY 1 AGTTCATCAAGACCGCCGATACGGCCACTTGGCCGTCAGACGCCGACCTTCACCTGC 60  
 Db 361 ArgPheLeuLysThrAlaIatyrGlyHisPheGlyArgAspAspProAspPheThrTrp 380

OY 61 GAGGTGTCACAGCC 75  
 Db 381 GluValValIysPro 385

RESULT 5  
 S46538  
 methionine adenosyltransferase (EC 2.5.1.6) - tomato  
 N:Alternate names: S-adenosyl-L-methionine synthetase  
 C:Species: Lycopersicon esculentum (tomato)  
 C:Date: 26-Dec-1994 #sequence\_revision 10-Nov-1995 #text\_change 05-May-2000  
 C:Accession: S46538; S38874  
 R:Espartero, J.; Pintor-Toro, J.A.; Pardo, J.M.  
 Plant Mol. Biol. 25, 217-227, 1994  
 A:Title: Differential accumulation of S-adenosylmethionine synthetase transcripts in  
 A:Reference number: S46538; MUID:94289646; PMID:8018871  
 A:Accession: S46538  
 A:Status: nucleic acid sequence not shown  
 A:Molecule type: mRNA  
 A:Residues: 1-393 <ES2>  
 A:Cross-references: EMBL:Z24741; NID:q429103; PIDN:CAA80865.1; PID:q429104  
 C:Superfamily: methionine adenosyltransferase  
 C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
 Pred. No.: 1.31e-09 Length: 393  
 Score: 120.00 Matches: 22  
 Percent Similarity: 92.00% Conservative: 1  
 Best Local Similarity: 88.00% Mismatches: 2  
 Query Match: 86.96% Indels: 0  
 DB: 2 Gaps: 0

US-09-198-779B-1\_COPY\_160\_235 (1-76) x S46538 (1-393)



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Alignment Scores:
Pred. No.: 7.38e-09 Length: 390
Score: 115.00 Matches: 22
Percent Similarity: 88.00% Conservative: 0
Best Local Similarity: 88.00% Mismatches: 3
Query Match: 83.33% Indels: 0
Gaps: 0

US-09-198-779B-1_COPY_160_235 (1-76) x G64785 (1-390)

OY 1 AGGTTTCATCAGACCGCGCATACGCCACTTGGCCGTGACGACCGCAGCTTCACTTC 60
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 361 ArgPheGlnLysThrAlaAlaLysrGlHisPheGlyArgAspAspProAspPheThrTrp 380
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

OY 61 GAGGTGTCACAGCCC 75
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 381 GluValValLysPro 385

RESULT 10
J00410
methionine adenosyltransferase (EC 2.5.1.6) 2 - Arabidopsis thaliana
N:Alternate names: S-adenosylmethionine synthetase
C:Species: Arabidopsis thaliana (mouse-ear cress)
C:Date: 12-Feb-1993 #sequence_revision 12-Feb-1993 #text_change 05-May-2000
A:Accession: J00410
R:Peleman, J.; Salto, K.; Cottyn, B.; Engler, G.; Seurinck, J.; Van Montagu, M.; Inze, L.
Gene 84, 359-369, 1989
A:Title: Structure and expression analyses of the S-adenosylmethionine synthetase gene
A:Reference number: J00410; MUID:90128280; PMID:2482229
A:Accession: J00410
A:Molecule type: DNA
A:Residues: 1-393 <PEL>
A:Cross-references: GB:M33217; NID:g166873; PIDN:AAA32869.1; PID:g166874
C:Genetics:
A:Gene: sam-2
C:Superfamily: methionine adenosyltransferase
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:
Pred. NO.: 7.38e-09 Length: 393
Score: 115.00 Matches: 22
Percent Similarity: 88.00% Conservative: 0
Best Local Similarity: 88.00% Mismatches: 3
Query Match: 83.33% Indels: 0
Gaps: 0

US-09-198-779B-1_COPY_160_235 (1-76) x J00410 (1-393)

OY 1 AGGTTTCATCAGACCGCGCATACGCCACTTGGCCGTGACGACCGCAGCTTCACTTC 60
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 361 ArgPheGlnLysThrAlaAlaLysrGlHisPheGlyArgAspAspProAspPheThrTrp 380
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

OY 61 GAGGTGTCACAGCCC 75
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 381 GluValValLysPro 385

RESULT 11
S66352
methionine adenosyltransferase (EC 2.5.1.6) 2 - garden pea
C:Species: Pisum sativum (garden pea)
C:Date: 14-Feb-1997 #sequence_revision 13-Mar-1997 #text_change 05-May-2000
A:Accession: S66352; S52218
R:Gomez-Gomez, L.; Carrasco, P.
Plant Mol. Biol. 30, 821-832, 1996
A:Title: Hormonal regulation of S-adenosylmethionine synthase transcripts in pea ovaries
A:Reference number: S66351; MUID:96194463; PMID:8624412
A:Accession: S66352
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-397 <GOM>
A:Cross-references: EMBL:X82077
R:Gomez, L.; Carrasco, P.

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submitted to the EMBL Data Library, October 1994
A:Description: Hormonal regulation of the s-adenosylmethionine synthase in pea ovarie
A:Reference number: S52218
A:Accession: S52218
A>Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-364,'FEDSCITWTFW' <GOW>
A:Cross-references: EMBL:X82077; NID:g609224; PIDN:CAA57581.1; PID:g609225
C:Superfamily: methionine adenosyltransferase
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:
Pred. No.:          7e-08                      Length:      397
Score:              108.50                     Matches:     22
Percent Similarity: 88.46%                     Conservative: 1
Best Local Similarity: 84.62%                   Mismatches:  2
Query Match:        78.62%                      Indels:      1
DB:                  2                          Gaps:        1

US-09-198-779B-1_COPY_160_235 (1-76) x S66352 (1-397)
Oy       1 AGGTTC--ATCAAGACGGCGCATACGCCACTTGGCCCTGCAGACGCCACTTCACC 57
         ||||| :||||:||||:||||:||||:||||:||||:||||:||||:||||:||||
Db       363 ARGPhelLeuleUlvSThAlAlAlatYrclYlnSpheGlYserhAspSPAlAAsPneThr 382
Oy       58 TGCAGGTGTCACGCC 75
         ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db       383 TrpGIuValIalYsPro 388

RESULT 12
707899
methionine adenosyltransferase (EC 2.5.1.6) - Chlamydomonas reinhardtii (fragment)
N:Alternate names: S-adenosylmethionine synthetase
C:Species: Chlamydomonas reinhardtii
C>Date: 14-May-1999 #sequence_revision 14-May-1999 #text_change 05-May-2000
C:Accession: T07899
R:Kim, J.Y.; Lee, K.O.; Lee, S.H.
Submitted to the EMBL Data Library, June 1997
A:Description: Chlamydomonas reinhardtii mRNA for S-adenosylmethionine synthetase.
A:Reference number: Z16198
A:Accession: T07899
A>Status: preliminary; translated from GB/EMBL/DDBJ
A:Molecule type: mRNA
A:Residues: 1-179 <KIM>
A:Cross-references: EMBL:AF008568; NID:g2454483; PIDN:AAB71833.1; PID:g2454484
A:Experimental source: strain 137C
C:Genetics:
A:Gene: SAMS
C:Function:
A:Description: catalyzes the formation of S-adenosyl methionine wth phosphate and py
C:Superfamily: methionine adenosyltransferase
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:
Pred. No.:          9.4e-07                      Length:      179
Score:              101.00                     Matches:     19
Percent Similarity: 83.33%                     Conservative: 1
Best Local Similarity: 79.17%                   Mismatches:  4
Query Match:        73.19%                      Indels:      0
DB:                  2                          Gaps:        0

US-09-198-779B-1_COPY_160_235 (1-76) x T07899 (1-179)
Oy       1 AGGTTCAACAGACGGCGCATACGCCACTTGGCCGTGACGCGCGCACTTCACCTGC 60
         |||:::||||:||||:||||:||||:||||:||||:||||:||||:||||:||||
Db       153 ARgYrGLnLstHrAlAlAtYrclYlnSpheGlYArGspAsPrroAsPneThrTrp 172
Oy       61 GAGGTGTCAG 72
         ||| |||||
Db       173 GlutHrValYls 176

RESULT 13
749491
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Best Local Similarity: 66.67%      Mismatches: 5
Query Match: 47.10%      Indels: 0
DB: 2      Gaps: 0

US-09-198-779B-1_COPY_160_235 (1-76) x D65657 (1-400)

QY 10 AAGACCGCGCATPACGGCCACTTTGGCGGTGACGACGGCGACTTCACCTGCGAG 63
DB 368 GlnThrAlaIatArgIyHisPheGlyArgHisAspValaAspLeuProTrpIu 385

RESULT 21
A:adenosylmethionine synthetase U0412 [Imported] - Ureaplasma urealyticum
C:Species: Ureaplasma urealyticum
C:Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 02-Sep-2000
C:Accession: A82895
R:Glass, J.I.; Letkowitz, E.J.; Glass, J.S.; Heiner, C.R.; Chen, E.Y.; Cassell, G.H.
submitted to Genbank, February 2000
A:Description: The complete sequence of Ureaplasma urealyticum: Alternate views of a mit
A:Reference number: A82870
A:Accession: A82895
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-376 <GLA>
A:Cross-references: GB:AE002138; GB:AF222894; NID:96899390; PIDN:AAF30823.1; GSPDB:GN001
A:Experimental source: serovar 3; biovar 1
C:Genetics:
A:Gene: mek; U0412
A:Genetic code: SGC3
C:Superfamily: methionine adenosyltransferase

Alignment Scores:
Pred. No.: 0.342      Length: 376
Score: 64.00      Matches: 10
Percent Similarity: 71.43%      Conservative: 5
Best Local Similarity: 47.62%      Mismatches: 6
Query Match: 46.38%      Indels: 0
DB: 2      Gaps: 0

US-09-198-779B-1_COPY_160_235 (1-76) x A82895 (1-376)

QY 1 AGGTCATCAGACAGCGCGCATACGGCCACTTGGCGGTGACGACGGCGACTTCACCTGC 60
DB 343 LysTyrLeuProValaIatThrTyrGlyHisPheGlyArgAspAspLeuAsnLeuSerTrp 362

QY 61 GAG 63
DB 363 Glu 363

RESULT 22
S27257
methionine adenosyltransferase (EC 2.5.1.6) 2 alpha chain - human
N:Alternate names: renal methionine adenosyltransferase (MAT); S-adenosylmethionine synt
C:Species: Homo sapiens (man)
C:Date: 22-Nov-1993 #sequence_revision 10-Nov-1995 #text_change 11-Jun-1999
C:Accession: S27257
R:Horikawa, S.; Tsukada, K.
FEBS Lett. 312, 37-41, 1992
A:Title: Molecular cloning and developmental expression of a human kidney S-adenosylmeth
A:Reference number: S27257; MUID:93050159; PMID:1426236
A:Accession: S27257
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-395 <HOR>
A:Cross-references: EMBL:X68836; GB:SA7859; NID:936326; PIDN:CAA48726.1; PID:936327
C:Genetics:
A:Gene: GDB:MAT2A; SAMS2; MAT2A
A:Cross-references: GDB:I36213; OMIM:601468
A:Map position: 2p11.2-2p11.2
A:Introns: 15/2; 256/3
C:Complex: heterodimer of catalytic alpha and regulatory beta chains
C:Function:
A:Description: catalyzes the formation of S-adenosyl methionine with phosphate and pyrop

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A:Pathway: one-carbon metabolism
C:Superfamily: methionine adenosyltransferase
C:Keywords: ATP; heterodimer; kidney; magnesium; metalloprotein; one-carbon metabolism
F:276-286/Region: nucleotide-binding motif A (P-loop) #status atypical
F:31/Binding site: magnesium 2 (Asp) #status predicted
F:285,289/Active site: Lys #status predicted
F:291/Binding site: magnesium 1 (Asp) #status predicted

Alignment Scores:
Pred. No.: 0.483      Length: 395
Score: 63.00      Matches: 14
Percent Similarity: 71.43%      Conservative: 1
Best Local Similarity: 66.67%      Mismatches: 4
Query Match: 45.65%      Indels: 2
DB: 1      Gaps: 1

US-09-198-779B-1_COPY_160_235 (1-76) x S27257 (1-395)

QY 10 AAGACCGCGCATPACGGCCACTTTGGCGGTGACGACGGCGACTTCACCTGCGAGGTGTC 69
DB 373 ArgThrAlaIatArgIyHisPheGlyArgAsp-----SerPheProTrpIuValaPro 390

QY 70 ANG 72
DB 391 Lys 391

RESULT 23
A37118
methionine adenosyltransferase (EC 2.5.1.6) - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 15-Feb-1991 #sequence_revision 15-Feb-1991 #text_change 05-May-2000
C:Accession: A37118
R:Horikawa, S.; Sasuga, J.; Shimizu, K.; Ozasa, H.; Tsukada, K.
J. Biol. Chem. 265, 13683-13686, 1990
A:Title: Molecular cloning and nucleotide sequence of cDNA encoding the rat kidney S-
A:Reference number: A37118; MUID:90337979; PMID:1696256
A:Accession: A37118
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-395 <HOR>
A:Cross-references: GB:J05571; NID:9206845; PIDN:AAA42106.1; PID:9206846
C:Superfamily: methionine adenosyltransferase
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:
Pred. No.: 0.483      Length: 395
Score: 63.00      Matches: 14
Percent Similarity: 71.43%      Conservative: 1
Best Local Similarity: 66.67%      Mismatches: 4
Query Match: 45.65%      Indels: 2
DB: 1      Gaps: 1

US-09-198-779B-1_COPY_160_235 (1-76) x A37118 (1-395)

QY 10 AAGACCGCGCATPACGGCCACTTTGGCGGTGACGACGGCGACTTCACCTGCGAGGTGTC 69
DB 373 ArgThrAlaIatArgIyHisPheGlyArgAsp-----SerPheProTrpIuValaPro 390

QY 70 ANG 72
DB 391 Lys 391

RESULT 24
H86976
probable S-adenosylmethionine synthase [Imported] - Mycobacterium leprae
C:Species: Mycobacterium leprae
C:Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 10-May-2001
C:Accession: H86976
R:Coile, S.T.; Eigmeier, K.; Parkhill, J.; James, K.D.; Thomson, N.R.; Wheeler, P.R.;
R.; Davies, R.M.; Devlin, K.; Duthey, S.; Feltwell, T.; Fraser, A.; Hamlin, N.; Holro
eam, M.A.; Rutherford, K.M.
Nature 409, 1007-1011, 2001
A:Authors: Rutter, S.; Seeger, K.; Simon, S.; Simmonds, M.; Skelton, J.; Squares, R.;

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Nature 397, 176-180, 1999  
A>Title: Genomic sequence comparison of two unrelated isolates of the human gastric p  
A:Reference number: A71800; MUID:99120557; PMID:9923682  
A:Accession: D71964  
A>Status: Preliminary  
A:Molecule type: DNA  
A:Residues: 1-385 <ARN>  
A:Cross-references: GB:AE001456; GB:AE001439; NID:94154689; PIDN:AAD05755.1; PID:9415  
A:Experimental source: strain J99  
C:Genetics:  
A:Gene: mekK  
C:Superfamily: methionine adenosyltransferase

Alignment Scores:

Pred. No.:	Length:	Matches:
0.965	385	11
Score:	61.00	Conservative: 3
Percent Similarity:	82.35%	Mismatches: 3
Best Local Similarity:	64.71%	Indels: 0
Query Match:	44.20%	Gaps: 0

DB: 2

US-09-198-779B-1.COPY\_160\_235 (1-76) x D71964 (1-385)

OY 13 ACCGGCGATACGCGCACTTGGCCGTGCAGACGCCGACTTCACCTGCGAG 63  
|||||:|||||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||  
Db 355 ThrselatatyrglyhisphneglYarggluenglugluinuphetrttppglu 371

RESULT 27 E64544

methionine adenosyltransferase (EC 2.5.1.6) 2 - Helicobacter pylori (strain 2695)  
C:Species: Helicobacter pylori  
C>Date: 09-Aug-1997 #sequence\_revision 09-Aug-1997 #text\_change 05-May-2000  
C:Accession: E64544  
R:Tomb, J.F.; White, O.; Kerlavage, A.R.; Clayton, R.A.; Sutton, G.G.; Fleischmann, R.  
Peterson, S.; Loftus, B.; Richardson, D.; Dodson, R.; Khalek, H.G.; Glodek, A.; McKee  
son, J.D.; Kelley, J.M.; Cotton, M.D.; Weidman, J.M.; Fujii, C.; Bowman, C.; Wathey,  
Nature 388, 539-547, 1997  
A:Authors: Wallin, E.; Hayes, W.S.; Borodovsky, M.; Karpk, P.D.; Smith, H.O.; Fraser,  
A>Title: The complete genome sequence of the gastric pathogen Helicobacter pylori.  
A:Reference number: A64520; MUID:97394467; PMID:9252185  
A:Accession: E64544  
A>Status: preliminary; nucleic acid sequence not shown; translation not shown  
A:Molecule type: DNA  
A:Residues: 1-385 <TOM>  
A:Cross-references: GB:AE000540; GB:AE000511; NID:92313287; PIDN:AAD07267.1; PID:9231  
C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:

Pred. No.:	Length:	Matches:
0.965	385	11
Score:	61.00	Conservative: 3
Percent Similarity:	82.35%	Mismatches: 3
Best Local Similarity:	64.71%	Indels: 0
Query Match:	44.20%	Gaps: 0

DB: 2

US-09-198-779B-1.COPY\_160\_235 (1-76) x E64544 (1-385)

OY 13 ACCGGCGATACGCGCACTTGGCCGTGCAGACGCCGACTTCACCTGCGAG 63  
|||||:|||||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||  
Db 355 ThrselatatyrglyhisphneglYarggluenglugluinuphetrttppglu 371

RESULT 28 F89964

S-adenosylmethionine synthetase [imported] - Staphylococcus aureus (strain N315)  
C:Species: Staphylococcus aureus  
C>Date: 10-May-2001 #sequence\_revision 10-May-2001 #text\_change 22-Oct-2001  
C:Accession: F89964  
R:Kuroda, M.; Ohita, Y.; Uchiyama, I.; Baba, T.; Yuzawa, H.; Kobayashi, I.; Cui, L.; O  
ma, A.; Mizutani-Ui, Y.; Kobayashi, N.; Sawano, T.; Inoue, R.; Kaito, C.; Sekimizu, K  
C.: Shiba, T.; Hattori, M.; Ogasawara, N.; Hayashi, H.; Hiramatsu, K.  
J: Nature 357, 1225-1240, 2001  
A>Title: Whole genome sequencing of methicillin-resistant Staphylococcus aureus.

A:Reference number: A89758; MUID:21311952; PMID:11418146  
A:Accession: F89964  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-398 <KIR>  
A:Cross-references: GB:BA000018; PID:913701583; PIDN:BA842876.1; GSPDB:GN00149  
A:Experimental source: strain N315  
C:Genetics:  
A:Gene: metK  
C:Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 0.965 Length: 398  
Score: 61.00 Matches: 12  
Percent Similarity: 72.22% Conservative: 1  
Best Local Similarity: 66.67% Mismatches: 5  
Query Match: 44.20% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779b-1\_COPY\_160\_235 (1-76) x F89964 (1-398)

OY 10 AAGACCGCGCATACGGCCACTTGGCCGTGACGACGCCGACTTCACCTGCGAG 63  
Db 367 GlnThrAlaAlaIatYrGlyHisPheGlyArGThrAspValaIGluLeuProTprGlu 384

RESULT 29  
D84062  
S:adenosylmethionine synthetase metK [imported] - Bacillus halodurans (strain C-125)  
C:Species: Bacillus halodurans  
C:Date: 01-Dec-2000 #sequence\_revision 01-Dec-2000 #text\_change 15-Jun-2001  
C:Accession: D84062  
R:Takami, H.; Nakasone, K.; Takaki, Y.; Maeno, G.; Sasaki, R.; Masui, N.; Fujii, F.; Hirai  
Nucleic Acids Res. 28, 4317-4331, 2000  
A:Title: Complete genome sequence of the alkaliphilic bacterium Bacillus halodurans and  
A:Reference number: A83650; MUID:20512582; PMID:11058132  
A:Accession: D84062  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-399 <STO>  
A:Cross-references: GB:AP001518; GB:BA000004; NID:910175792; PIDN:BA807019.1; GSPDB:GN00  
C:Genetics:  
A:Experimental source: strain C-125  
C:Genetics:  
A:Gene: metK  
C:Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 0.965 Length: 399  
Score: 61.00 Matches: 11  
Percent Similarity: 72.22% Conservative: 2  
Best Local Similarity: 61.11% Mismatches: 5  
Query Match: 44.20% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779b-1\_COPY\_160\_235 (1-76) x D84062 (1-399)

OY 10 AAGACCGCGCATACGGCCACTTGGCCGTGACGACGCCGACTTCACCTGCGAG 63  
Db 370 GlnThrAlaAlaIatYrGlyHisPheGlyArGThrAspValaIGluLeuProTprGlu 387

RESULT 30  
F86862  
methionine adenosyltransferase (EC 2.5.1.6) [imported] - Lactococcus lactis subsp. lacti  
N:Alternate names: S-adenosylmethionine synthetase  
C:Species: Lactococcus lactis subsp. lactis  
C:Date: 23-Mar-2001 #sequence\_revision 23-Mar-2001 #text\_change 03-Aug-2001  
C:Accession: F86862  
R:Boilot, A.; Wincker, P.; Mauger, S.; Jallion, O.; Malarne, K.; Weissenbach, J.; Ehrlich  
Genome Res. 11, 731-753, 2001  
A:Title: The complete genome sequence of the lactic acid bacterium Lactococcus lactis ss  
A:Reference number: A86625; MUID:21235186; PMID:11337471  
A:Accession: F86862  
A:Status: preliminary  
A:Molecule type: DNA

A:Residues: 1-399 <STO>  
A:Cross-references: GB:AE005176; PID:912724937; PIDN:AAK06000.1; GSPDB:GN00146  
A:Experimental source: strain IL1403  
C:Genetics:  
A:Gene: metK  
C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
Pred. No.: 0.965 Length: 399  
Score: 61.00 Matches: 11  
Percent Similarity: 72.22% Conservative: 2  
Best Local Similarity: 61.11% Mismatches: 5  
Query Match: 44.20% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779b-1\_COPY\_160\_235 (1-76) x F86862 (1-399)

OY 10 AAGACCGCGCATACGGCCACTTGGCCGTGACGACGCCGACTTCACCTGCGAG 63  
Db 368 GlnThrAlaAlaIatYrGlyHisPheGlyArGThrAspValaIGluLeuProTprGlu 385

RESULT 31  
AD1654  
S:methionine adenosyltransferase homolog metK [imported] - Listeria innocua (strain C  
C:Species: Listeria innocua  
C:Date: 27-Nov-2001 #sequence\_revision 27-Nov-2001 #text\_change 14-Dec-2001  
C:Accession: AD1654  
R:Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloec  
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entlian, K.D.; Fsihl,  
D.; Jones, L.M.; Karst, U.  
Science 294, 849-852, 2001  
A:Authors: Kretz, J.; Kuhn, M.; Kunst, F.; Kurapkay, G.; Madueno, E.; Maltournam, A.;  
Ok, C.; Schueter, T.; Simoes, N.; Tilleret, A.; Vazquez-Boland, J.A.; Voss, H.; Wehla  
A:Title: Comparative genomics of Listeria species  
A:Reference number: AB1077; MUID:21537279; PMID:11679669  
A:Accession: AD1654  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-399 <GLA>  
A:Cross-references: GB:AL592022; PIDN:CACG7004.1; PID:916414260; GSPDB:GN00178  
A:Experimental source: strain Clp11262  
C:Genetics:  
A:Gene: metK  
C:Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 0.965 Length: 399  
Score: 61.00 Matches: 11  
Percent Similarity: 72.22% Conservative: 2  
Best Local Similarity: 61.11% Mismatches: 5  
Query Match: 44.20% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779b-1\_COPY\_160\_235 (1-76) x AD1654 (1-399)

OY 10 AAGACCGCGCATACGGCCACTTGGCCGTGACGACGCCGACTTCACCTGCGAG 63  
Db 368 GlnThrAlaAlaIatYrGlyHisPheGlyArGThrAspValaIGluLeuProTprGlu 385

RESULT 32  
AH1282  
S:methionine adenosyltransferase homolog metK [imported] - Listeria monocytogenes (st  
C:Species: Listeria monocytogenes  
C:Date: 27-Nov-2001 #sequence\_revision 27-Nov-2001 #text\_change 14-Dec-2001  
C:Accession: AH1282  
R:Glaser, P.; Frangeul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloec  
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entlian, K.D.; Fsihl,  
D.; Jones, L.M.; Karst, U.  
Science 294, 849-852, 2001  
A:Authors: Kretz, J.; Kuhn, M.; Kunst, F.; Kurapkay, G.; Madueno, E.; Maltournam, A.;  
Ok, C.; Schueter, T.; Simoes, N.; Tilleret, A.; Vazquez-Boland, J.A.; Voss, H.; Wehla  
A:Title: Comparative genomics of Listeria species.

A:Reference number: AB1077; MUID:21537279; PMID:11679666  
A:Accession: AH1282  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-399 <GIA>  
A:Cross-references: GB:NC\_003210; PIDN:CAC99742.1; PID:g16411100; GSPDB:GN00177  
A:Experimental source: strain ESD-e  
C:Genetics:  
A:Gene: metK  
C:Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 0.965 Length: 399  
Score: 61.00 Matches: 11  
Percent Similarity: 72.22% Conservative: 2  
Best Local Similarity: 61.11% Mismatches: 5  
Query Match: 44.20% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779b-1\_COPY\_160\_235 (1-76) x AH1282 (1-399)

OY 10 AAGACCGCCGATACGCGCACTTGGCGTGAAGCGCGACTTCACCTGCGAG 63  
::: |||||::: ||||| ||| ||| |||  
DB: 368 GlnThrAlaIalalaphcglYhlsphcglYargserAspLeuAspLeuProtrpGlu 385

RESULT 33  
F70899  
Probable metK protein - Mycobacterium tuberculosis (strain H37RV)  
C:Species: Mycobacterium tuberculosis  
C>Date: 17-Jul-1998 #sequence\_revision 17-Jul-1998 #text\_change 20-Jun-2000  
C:Accession: F70899  
R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.; Connor, R.; Davies, R.; Devlin, K.; Fellwell, T.; Genovesi, S.; Hamlin, N.; Holroyd, S.; Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.  
Nature 393, 537-544, 1998  
A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.  
A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete genome  
A:Reference number: A70500; MUID:9829587; PMID:9634230  
A:Accession: F70899  
A:Status: preliminary; nucleic acid sequence not shown; translation not shown  
A:Molecule type: DNA  
A:Residues: 1-403 <COL>  
A:Cross-references: GB:Z80108; GB:AL123456; NID:33256012; PIDN:CAN02194.1; PID:g3356013  
A:Experimental source: strain H37RV  
C:Genetics:  
A:Gene: metK  
C:Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 0.965 Length: 403  
Score: 61.00 Matches: 11  
Percent Similarity: 65.00% Conservative: 2  
Best Local Similarity: 55.00% Mismatches: 7  
Query Match: 44.20% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779b-1\_COPY\_160\_235 (1-76) x F70899 (1-403)

OY 4 TTCATCAAGACCGCGCATACGCGCACTTGGCGTGAAGCGACTTCACCTGCGAG 63  
::: |||||::: ||||| ||| ||| |||  
DB: 372 TyrAlaProThrAlaIalalYrglYhlsphcglYargThrAspValcIuLeuProtrpGlu 391

RESULT 34  
B97403  
methionine adenosyltransferase (EC 2.5.1.6) - Agrobacterium tumefaciens (strain C58, C58, C58)  
C:Species: Agrobacterium tumefaciens  
C>Date: 30-Sep-2001 #sequence\_revision 30-Sep-2001 #text\_change 11-Jan-2002  
C:Accession: B97403  
R:Goodner, B.; Hinkle, G.; Gelling, S.; Miller, N.; Blanchard, M.; Qurollo, B.; Goldman, A.; Liu, F.; Wollam, C.; Allinger, M.; Doughty, D.; Scott, C.; Lappas, C.; Markelz, B.; Science 294, 2333-2328, 2001  
A:Title: Genome Sequence of the Plant Pathogen and Biotechnology Agent Agrobacterium tumefaciens  
A:Reference number: A97359; PMID:11743194

A:Accession: B97403  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-420 <KUR>  
A:Cross-references: GB:AE007869; PIDN:AAK86179.1; PID:g15155274; GSPDB:GN00169  
C:Genetics:  
A:Gene: AGR\_C\_632  
A:Map position: circular chromosome  
C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
Pred. No.: 0.965 Length: 420  
Score: 61.00 Matches: 14  
Percent Similarity: 63.33% Conservative: 5  
Best Local Similarity: 46.67% Mismatches: 5  
Query Match: 44.20% Indels: 6  
DB: 2 Gaps: 2

US-09-198-779b-1\_COPY\_160\_235 (1-76) x B97403 (1-420)

OY 4 TTCATCAAGACCGCGCATACGCGCACTTGGCGTGAAC-----GACGCGGACTTC 54  
::: |||||::: ||||| ||| ||| |||  
DB: 384 TyrAlaIalYhlsphcglYhlsphcglYargYsalagYargAspGlySerPhe 403

OY 55 ACCTGCGAG-----GTGGTCAAGCCG 75  
::: |||  
DB: 404 SerTrpGluYlsLeuAspLeuValYlsPro 413

RESULT 35  
AB2621  
S-adenosylmethionine synthetase metK [imported] - Agrobacterium tumefaciens (strain C58)  
C:Species: Agrobacterium tumefaciens  
C>Date: 11-Jan-2002 #sequence\_revision 11-Jan-2002 #text\_change 01-Feb-2002  
C:Accession: AB2621  
R:Wood, D.W.; Setubal, J.C.; Kaul, R.; Monks, D.; Chen, L.; Wood, G.E.; Chen, Y.; Woc-erage, G.; Gillet, M.; Grant, C.; Guenther, D.; Kutayvin, T.; Levy, R.; Li, M.; McCl ; Karp, P.; Romero, P.; Zhang, S.  
Science 294, 2317-2323, 2001  
A:Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry, M.; Gordon-Kamster, E.W.  
A:Title: The genome of the Natural Genetic Engineer Agrobacterium tumefaciens C58.  
A:Reference number: AB2577; PMID:11743193  
A:Accession: AB2621  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-420 <KUR>  
A:Cross-references: GB:AE008688; PIDN:AA141384.1; PID:g17738701; GSPDB:GN00186  
A:Experimental source: strain C58 (Dupont)  
C:Genetics:  
A:Gene: metK  
A:Map position: circular chromosome  
C:Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 0.965 Length: 420  
Score: 61.00 Matches: 14  
Percent Similarity: 63.33% Conservative: 5  
Best Local Similarity: 46.67% Mismatches: 5  
Query Match: 44.20% Indels: 6  
DB: 2 Gaps: 2

US-09-198-779b-1\_COPY\_160\_235 (1-76) x AB2621 (1-420)

OY 4 TTCATCAAGACCGCGCATACGCGCACTTGGCGTGAAC-----GACGCGGACTTC 54  
::: |||||::: ||||| ||| ||| |||  
DB: 384 TyrAlaIalYhlsphcglYhlsphcglYargYsalagYargAspGlySerPhe 403

OY 55 ACCTGCGAG-----GTGGTCAAGCCG 75  
::: |||  
DB: 404 SerTrpGluYlsLeuAspLeuValYlsPro 413

RESULT 36

S74736  
 methionine adenosyltransferase (EC 2.5.1.6) - *Synechocystis* sp. (strain PCC 6803)  
 N:Alternate names: protein sll0927; S-adenosylmethionine synthetase  
 C:Species: *Synechocystis* sp.  
 A:Variety: PCC 6803  
 C:Date: 25-Apr-1997 #sequence, revision 25-Apr-1997 #text\_change 20-Jun-2000  
 C:Accession: S74736  
 R:Kaneko, T.; Sato, S.; Kotani, H.; Tanaka, A.; Asamizu, E.; Nakamura, Y.; Miyajima, N.; O. K.; Okumura, S.; Shimo, S.; Takeuchi, C.; Wada, T.; Matsubae, A.; Yamada, M.; Yasuda  
 DNA Res. 3: 109-136, 1996  
 A:Title: Sequence analysis of the genome of the unicellular cyanobacterium *Synechocystis*  
 S.  
 A:Reference number: S74322; MUID:97061201; PMID:8905231  
 A:Accession: S74736  
 A:Status: nucleic acid sequence not shown; translation not shown  
 A:Molecule type: DNA  
 A:Residues: 1-409 <KAN>  
 A:Cross-references: EMBL:D90901; GB:AE001339; NID:91651897; PIDN:BA16887.1; PID:9165196  
 A:Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1996  
 C:Genetics:  
 A:Gene: metX  
 A:Start codon: GTG  
 C:Superfamily: methionine adenosyltransferase  
 C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
 Pred. No.: 1.15 Length: 409  
 Score: 60.50 Matches: 13  
 Percent Similarity: 68.18% Conservative: 2  
 Best Local Similarity: 59.09% Mismatches: 6  
 Query Match: 43.84% Indels: 1  
 Gaps: 2

US-09-198-779b-1\_COPY\_160\_235 (1-76) x S74736 (1-409)

OY 1 AGGTCATCAG---ACCGCCGATACGGCCACTTGGCCGTGACGACCCGACTTACAC 57  
 Db 367 ArgPterYrGlnAspValAlaIatYrGlyHisPheGlyArgAsnAspLeuAspLeuPro 386

OY 58 TGGCAG 63  
 Db 387 TrpGlu 388

RESULT 37  
 E82319  
 S-adenosylmethionine synthase WC0472 [imported] - *Vibrio cholerae* (strain N16961 serogroup  
 C:Species: *Vibrio cholerae*  
 C:Date: 18-Aug-2000 #sequence, revision 20-Aug-2000 #text\_change 02-Feb-2001  
 C:Accession: E82319  
 R:Heidelberg, J.F.; Eisen, J.A.; Nelson, W.C.; Clayton, R.A.; Gwinn, M.L.; Dodson, R.J.;  
 charlson, D.; Ermolaeva, M.D.; Vamathevan, J.; Bass, S.; Qin, H.; Dragol, I.; Sellers, F.  
 L.; R.R.; Mekalanos, J.J.; Venter, J.C.; Fraser, C.M.  
 Nature 406: 477-483, 2000  
 A:Title: DNA Sequence of both chromosomes of the cholera pathogen *Vibrio cholerae*.  
 A:Reference number: A82035; MUID:20406833; PMID:10952301  
 A:Accession: E82319  
 A:Status: preliminary  
 A:Molecule type: DNA  
 A:Residues: 1-385 <HEI>  
 A:Cross-references: GB:AE004133; GB:AE003852; NID:99654889; PIDN:AA93645.1; GSPDB:GN001  
 A:Experimental source: serogroup O1; strain N16961; biotype El Tor  
 C:Genetics:  
 A:Gene: VC0472  
 A:Map position: 1  
 C:Superfamily: methionine adenosyltransferase

Alignment Scores:  
 Pred. No.: 1.36 Length: 385  
 Score: 60.00 Matches: 10  
 Percent Similarity: 100.00% Conservative: 2  
 Best Local Similarity: 83.33% Mismatches: 0  
 Query Match: 43.48% Indels: 0  
 Gaps: 2

US-09-198-779b-1\_COPY\_160\_235 (1-76) x E82319 (1-385)

OY 10 AAGACCGCCGATACGGCCACTTGGCCGTGACGAC 45  
 Db 355 LysThrAlaIatYrGlyHisPheGlyArgGluGlu 366

RESULT 38  
 A71281  
 probable S-adenosylmethionine synthetase (metK) - *Syphilis spirochete*  
 C:Species: *Treponema pallidum* subsp. *pallidum* (*Syphilis spirochete*)  
 C:Date: 24-Jul-1998 #sequence, revision 24-Jul-1998 #text\_change 18-Jun-1999  
 C:Accession: A71281  
 R:Fraser, C.M.; Norris, S.J.; Weinstein, G.M.; White, O.; Sutton, G.G.; Dodson, R.; G  
 rson, J.; Khalak, H.; Richardson, D.; Howell, J.K.; Chidambaram, M.; Uitterback, T.; M  
 they, L.; Weidman, J.; Smith, H.O.; Venter, J.C.  
 Science 281: 375-388, 1998  
 A:Title: Complete genome sequence of *Treponema pallidum*, the *Syphilis spirochete*.  
 A:Reference number: A71250; MUID:98332770; PMID:9665876  
 A:Accession: A71281  
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown  
 A:Molecule type: DNA  
 A:Residues: 1-396 <COL>  
 A:Cross-references: GB:AE001250; GB:AE000520; NID:93323096; PIDN:AA65758.1; PID:9332  
 A:Experimental source: strain Nichols  
 C:Genetics:  
 A:Gene: TP0794  
 C:Superfamily: methionine adenosyltransferase

Alignment Scores:  
 Pred. No.: 1.93 Length: 396  
 Score: 59.00 Matches: 13  
 Percent Similarity: 54.55% Conservative: 5  
 Best Local Similarity: 39.39% Mismatches: 7  
 Query Match: 42.75% Indels: 8  
 Gaps: 2

US-09-198-779b-1\_COPY\_160\_235 (1-76) x A71281 (1-396)

OY 1 AGGTCATCAGACCGCCGATACGGCCACTTGGCCGTGACGAC----- 45.  
 Db 355 ArgYrArSerThrAlaIatYrGlyHisPheGlyArgGluGlnPheProTrpGluArg 374

OY 46 GCCGACTTCACCTGCGAGGTG-----GTCAAGCC 75  
 Db 375 ThrAspValCysAspLeuGlnArgAlaValArgPro 387

RESULT 39  
 A47151  
 methionine adenosyltransferase (EC 2.5.1.6) - mouse  
 N:Alternate names: S-adenosylmethionine synthetase  
 C:Species: *Mus musculus* (house mouse)  
 C:Date: 21-Jan-1994 #sequence, revision 18-Nov-1994 #text\_change 05-May-2000  
 C:Accession: A47151  
 R:Sakata, S.F.; Shelly, L.L.; Ruppert, S.; Schutz, G.; Chou, J.Y.  
 J. Biol. Chem. 268: 13978-13986, 1993  
 A:Title: Cloning and expression of murine S-adenosylmethionine synthetase.  
 A:Reference number: A47151; MUID:93300783; PMID:8314764  
 A:Accession: A47151  
 A:Status: preliminary  
 A:Molecule type: nucleic acid  
 A:Residues: 1-396 <SAK>  
 A:Experimental source: liver  
 A:Note: sequence inconsistent with the nucleotide translation;  
 C:Superfamily: methionine adenosyltransferase  
 C:Keywords: S-adenosylmethionine; transferase

Alignment Scores:  
 Pred. No.: 1.93 Length: 396  
 Score: 59.00 Matches: 13  
 Percent Similarity: 71.43% Conservative: 2  
 Best Local Similarity: 61.90% Mismatches: 4

Query Match: 42.75% Indels: 2  
DB: 2 Gaps: 1  
US-09-198-779B-1\_COPY\_160\_235 (1-76) x A47151 (1-396)  
QY 10 AAGACCGCCGATACGCGCCTTGGCCGTGACGACGCGGACTTCACCTCCGAGTGTC 69  
||||| ||||||| ||||||| ||||||| |||||||  
DB 374 LysThrAlaCysTyrGlyHisPheGlyArg-----SerGlnPheProTrrpGluValPro 391  
QY 70 AAG 72  
|||  
DB 392 Lys 392  
RESULT 40  
S06114  
methionine adenosyltransferase (EC 2.5.1.6) - rat  
N:Alternate names: S-adenosylmethionine synthetase  
C:Species: Rattus norvegicus (Norway rat)  
C:Date: 31-Mar-1990 #sequence\_revision 31-Mar-1990 #text\_change 05-May-2000  
C:Accession: S06114; S18256; S18257  
R:Horikawa, S.; Ishikawa, M.; Ozasa, H.; Tsukada, K.  
Eur. J. Biochem. 184, 497-501, 1989  
A:Title: Isolation of a cDNA encoding the rat liver S-adenosylmethionine synthetase.  
A:Reference number: S06114; MUID:90032633; PMID:2806235  
A:Accession: S06114  
A:Molecule type: mRNA  
A:Residues: 1-397 <HOR>  
A:Cross-references: EMBL:X15734; NID:957183; PIDN:CAA3754.1; PID:957184  
R:Mato, J.M.  
submitted to the EMBL Data Library, July 1991  
A:Reference number: S18256  
A:Accession: S18256  
A:Status: preliminary  
A:Molecule type: mRNA  
A:Residues: 1-397 <MAT>  
A:Cross-references: EMBL:X60822  
R:Alvarez, L.; Asuncion, M.; Corrales, F.; Pajares, M.A.; Mato, J.M.  
FEBS Lett. 290, 142-146, 1991  
A:Title: Analysis of the 5' non-coding region of rat liver S-adenosylmethionine synthetase  
A:Reference number: S18257; MUID:92008649; PMID:1915866  
A:Accession: S18257  
A:Status: translation not shown  
A:Molecule type: mRNA  
A:Residues: 1-46 <ALV>  
A:Cross-references: EMBL:X60822  
C:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase  
Alignment Scores:  
Pred. No.: 1.93 Length: 397  
Score: 59.00 Matches: 13  
Percent Similarity: 71.43% Conservative: 2  
Best Local Similarity: 61.90% Mismatches: 4  
Query Match: 42.75% Indels: 2  
DB: 2 Gaps: 1  
US-09-198-779B-1\_COPY\_160\_235 (1-76) x S06114 (1-397)  
QY 10 AAGACCGCCGATACGCGCCTTGGCCGTGACGACGCGGACTTCACCTCCGAGTGTC 69  
||||| ||||||| ||||||| ||||||| |||||||  
DB 375 LysThrAlaCysTyrGlyHisPheGlyArg-----SerGlnPheProTrrpGluValPro 392  
QY 70 AAG 72  
|||  
DB 393 Lys 393  
RESULT 41  
H70866  
hypothetical protein Rv2473 - Mycobacterium tuberculosis (strain H37RV)  
C:Species: Mycobacterium tuberculosis  
C:Date: 17-Jul-1998 #sequence\_revision 17-Jul-1998 #text\_change 22-Oct-1999  
C:Accession: H70866  
R:Cole, S.T.; Brosch, P.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.

; Connor, R.; Davies, R.; Devlin, K.; Fellwell, T.; Gentles, S.; Hamlin, N.; Holroyd, R.; Jandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.  
Nature 393, 537-544, 1998  
A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.  
A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete genome  
A:Reference number: A70500; MUID:98295987; PMID:9634230  
A:Accession: H70866  
A:Status: preliminary; nucleic acid sequence not shown; translation not shown  
A:Molecule type: DNA  
A:Residues: 1-238 <COL>  
A:Cross-references: GB:A1021246; GB:A1123456; NID:93261507; PIDN:CAA16050.1; PID:e123  
A:Experimental source: strain H37RV  
C:Genetics:  
A:Gene: Rv2473  
Alignment Scores:  
Pred. No.: 2.29 Length: 238  
Score: 58.50 Matches: 13  
Percent Similarity: 60.87% Conservative: 1  
Best Local Similarity: 56.52% Mismatches: 4  
Query Match: 42.39% Indels: 5  
DB: 2 Gaps: 1  
US-09-198-779B-1\_COPY\_160\_235 (1-76) x H70866 (1-238)  
QY 2 GGTTCATCAGACCGCGCATACGCGCCTTGGCCGTGACGACGCGGACTTCACCTGCG 61  
||||| ||||||| |||  
DB 66 GlySerSerArgProAlaProSer-----ThrThrProArgSerProGly 80  
QY 62 AGCTGCGCA 70  
|||||  
DB 81 ArgTrrSer 83  
RESULT 42  
E84977  
methionine adenosyltransferase (EC 2.5.1.6) [imported] - Buchnera sp. (strain APS)  
N:Alternate names: S-adenosylmethionine synthetase  
C:Species: Buchnera sp.  
C:Date: 02-Mar-2001 #sequence\_revision 02-Mar-2001 #text\_change 23-Mar-2001  
C:Accession: E84977  
R:Shigenobu, S.; Matanabe, H.; Hattori, M.; Sakaki, Y.; Ishikawa, H.  
Nature 407, 81-86, 2000  
A:Title: Genome sequence of the endocellular bacterial symbiont of aphids Buchnera sp.  
A:Reference number: A84930; MUID:20445173; PMID:10993077  
A:Accession: E84977  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-378 <STO>  
A:Cross-references: GB:AP000398; GSPDB:GM00144  
A:Experimental source: strain APS  
C:Genetics:  
A:Gene: melK; BU408  
A:Superfamily: methionine adenosyltransferase  
C:Keywords: S-adenosylmethionine; transferase  
Alignment Scores:  
Pred. No.: 2.73 Length: 378  
Score: 58.00 Matches: 9  
Percent Similarity: 85.71% Conservative: 3  
Best Local Similarity: 64.29% Mismatches: 2  
Query Match: 42.03% Indels: 0  
DB: 2 Gaps: 0  
US-09-198-779B-1\_COPY\_160\_235 (1-76) x E84977 (1-378)  
QY 4 TTCATCAAGACCGCGCATACGCGCCTTGGCCGTGACGAC 45  
||||| ||||||| |||  
DB 352 TyrLeuLysThrAlaValTyrGlyHisPheGlyArgGlySerGlu 365  
RESULT 43  
T13645  
microtubule-associated protein - fruit fly (Drosophila melanogaster)  
C:Species: Drosophila melanogaster



C>Date: 20-Sep-1999 #sequence\_revision 20-Sep-1999 #text\_change 17-Nov-2000  
C:Accession: T13845  
R:Saunders, R.D.; Avides, M.C.; Howard, T.; Gonzalez, C.; Glover, D.M.  
J. Cell Biol. 137, 881-890, 1997  
A:Title: The drosophila gene abnormal spindle encodes a microtubule-associated protein  
C:Accession: T13845  
A:Status: preliminary; translated from GB/EMBL/DBJ  
A:Molecule type: mRNA  
A:Residues: 1-1861 <SAU>  
A:Cross-references: EMBL:U95171; MID:91930121; PID:91930122; PIDN:AAB51540.1  
C:Genetics:  
A:Gene: asp  
A:Cross-references: Flybase:Fbgn0000140  
C:Function:  
A:Description: is required for the normal function of the mitotic spindle

Alignment Scores:  
Pred. No.: 2.72 Length: 1861  
Score: 58.00 Matches: 10  
Percent Similarity: 58.33% Conservative: 4  
Best Local Similarity: 41.67% Mismatches: 10  
Query Match: 42.03% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1\_COPY\_160\_235 (1-76) x T13845 (1-1861)

OY 5 TCATCAGAGCGCGCATACGCGCCACTTGGCCGCGAGCGAGCGACTTCACCTGCGAGG 64  
Db 83 AAlaAlaAlaProPseRerLysGlnThrTrpAlaValThrAlaProSerArgProAlaAla 102  
OY 65 TGCTCAAGCCCC 76  
Db 103 TrpAlaHisPro 106

RESULT 44  
AE2211  
S-adenosylmethionine synthetase [imported] - Nostoc sp. (strain PCC 7120)  
C:Species: Nostoc sp.  
A:Note: Nostoc sp. strain PCC 7120 is a synonym of Anabaena sp. strain PCC 7120  
C>Date: 14-Dec-2001 #sequence\_revision 14-Dec-2001 #text\_change 30-Jun-2002  
C:Accession: AE2211  
R:Kaneko, T.; Nakamura, Y.; Wolk, C.P.; Kuritz, T.; Sasamoto, S.; Watanabe, A.; Iriguchi, N.; Shimpo, S.; Sugimoto, M.; Takazawa, M.; Yasuda, M.; Tabata, S.  
DNA Res. 8, 205-213, 2001  
A:Title: Complete genomic sequence of the filamentous Nitrogen-fixing Cyanobacterium Anabaena  
A:Reference number: AB1807; MUID:21595285; PMID:11759840  
A:Accession: AE2211  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-394 <KUR>  
A:Cross-references: GB:BA000019; PIDN:BA874943.1; PID:917132339; GSPDB:GN00179  
C:Experimental source: strain PCC 7120  
C:Genetics:  
A:Gene: all3244  
C:Superfamily: methionine adenosyltransferase

Alignment Scores:  
Pred. No.: 3.85 Length: 394  
Score: 57.00 Matches: 11  
Percent Similarity: 60.00% Conservative: 1  
Best Local Similarity: 55.00% Mismatches: 8  
Query Match: 41.30% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1\_COPY\_160\_235 (1-76) x AE2211 (1-394)

OY 10 AAGACCGCGCATACGCGCACTTGGCCGCGAGCGAGCGACTTCACCTGCGAGGTGTC 69  
Db 369 LysLeuAlaAlaTyrGlyHisValGlyArgGlnAspLeuProTropIuLysIle 388

RESULT 45  
T50705

gamma-aminobutyrate aminotransferase (EC 2.6.1.-) rHba homolog [imported] - Rhodobact  
N:Alternate names: pyridoxal-phosphate dependent aminotransferase  
C:Species: Rhodobacter sphaeroides  
C>Date: 21-Jul-2000 #sequence\_revision 21-Jul-2000 #text\_change 21-Jul-2000  
C:Accession: T50705  
R:Choudhary, M.; Kaplan, S.  
Nucleic Acids Res. 28, 862-867, 2000  
A:Title: DNA sequence analysis of the photosynthesis region of Rhodobacter sphaeroides  
A:Reference number: Z25222; MUID:20115911; PMID:10648776  
A:Accession: T50705  
A:Status: preliminary; translated from GB/EMBL/DBJ  
A:Molecule type: DNA  
A:Residues: 1-447 <CHO>  
A:Cross-references: EMBL:AF195122; PIDN:AAF24249.1  
A:Experimental source: strain 2.4.1  
C:Genetics:  
A:Gene: rHba  
C:Keywords: aminotransferase

Alignment Scores:  
Pred. No.: 3.85 Length: 447  
Score: 57.00 Matches: 11  
Percent Similarity: 55.56% Conservative: 4  
Best Local Similarity: 40.74% Mismatches: 8  
Query Match: 41.30% Indels: 4  
DB: 2 Gaps: 1

US-09-198-779B-1\_COPY\_160\_235 (1-76) x T50705 (1-447)

OY 8 TCACAGCGCGCG-----CATACGCGCGCACTTGGCCGCGTACGAGCGCGCACTTCA 55  
Db 5 SerLeuProProArgGlyAlaAlaHisGluProAlaMetProCysArgThrProSerAla 24  
OY 56 CCTGCGAGGTGTCACGCCCC 76  
Db 25 ProAlaArgTyrThrCysPro 31

Search completed: April 23, 2003, 12:08:01  
Job time : 8.33333 secs

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[illegible]

## SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	121	100.0	2061	21	AAA48574	CDNA encoding corn
2	29.2	24.1	2604	21	AAA48576	CDNA encoding wheat
3	28.8	23.8	705	24	ABQ25086	Oligonucleotide for
4	28.8	23.8	705	24	ABQ25087	Oligonucleotide for
5	28.8	23.6	1519	20	AAV64373	GABA-gated chloride
6	28.6	23.6	405	23	ABV48454	Human prostate exp
7	28.2	23.3	297	16	AAQ8658	Beta-amyloid precu
8	28.2	23.3	300	24	AAK9387	DNA of wild type C
9	28	23.1	1357	21	AAC79021	Human secreted pried

Apr 23 15:43:09 2003

C	10	27.6	22.8	1086	23	ABL04431
C	11	27.6	22.8	3305	23	ABL04430
C	12	27.6	22.8	3457	23	ABL04426
C	13	27.4	22.6	4659	23	ABL29895
C	14	27.4	22.6	7643	23	ABL29894
C	15	27.4	22.6	32042	20	AAD209252
C	16	27.4	22.6	32042	22	AAE30011
C	17	27.4	22.6	32042	24	AAI40765
C	18	27	22.3	628	24	ABO25024
C	19	27	22.3	628	24	ABQ25025
C	20	26.8	22.1	574	23	ABV34453
C	21	26.8	22.1	408	22	AAI86054
C	22	26.8	22.1	486	21	AAZ297380
C	23	26.8	22.1	1004	21	AAZ297180
C	24	26.8	22.1	2033	22	AAD03458
C	25	26.8	22.1	2037	21	AAZ29961
C	26	26.8	22.1	2650	22	AAH14585
C	27	26.8	22.1	2837	22	AAH16527
C	28	26.8	22.1	3095	17	AAAT31290
C	29	26.8	22.1	13255	23	ABLI6634
C	30	26.8	22.1	29392	19	AAVL5422
C	31	26.8	22.1	33513	22	AAK78746
C	32	26.6	22.0	294	9	AAAN81965
C	33	26.6	22.0	297	16	AAO88696
C	34	26.6	22.0	297	16	AAQ88697
C	35	26.6	22.0	300	14	AAQ42665
C	36	26.6	22.0	300	24	AAK99386
C	37	26.6	22.0	303	21	AAA39488
C	38	26.6	22.0	309	16	AAO88699
C	39	26.6	22.0	309	16	AAO88700
C	40	26.6	22.0	315	22	AAD20982
C	41	26.6	22.0	327	24	AAD23935
C	42	26.6	22.0	336	17	AAAI8082
C	43	26.6	22.0	337	14	AAO59415
C	44	26.6	22.0	354	19	AAO7188
C	45	26.6	22.0	354	19	AAV20379
C	46	26.6	22.0	354	21	AAA39490
C	47	26.6	22.0	354	22	AAAD20981
C	48	26.6	22.0	495	21	AAA39492
C	49	26.6	22.0	581	22	AAAL15643
C	50	26.6	22.0	1062	9	AAAN80607

RESULT 1  
AAAA8574  
ID AAAA8574 standard; cdna; 2061 bp.

ALIGNMENTS

corn

CDNA encoding corn protein phosphatase 2A regulatory subunit A.  
Corn: protein phosphatase 2A; protein phosphorylation modulation;  
transgenic plant; gene therapy; ss.

Zea mays.

Key Location/Qualifiers  
CDS 56..1820 /+tag= a  
FT FT CDS /product= "protein phosphatase 2A regulatory  
FT FT subunit A"

WO200036121-A2.  
22-JUN-2000.  
15-DEC-1999; 99MO-US29823.

PR 16-DEC-1998; 98US-0112541.  
XX  
PA (DUPO ) DU PONT DE NEMOURS & CO E I.  
XX  
PI Famodu OO, Miao G, Sakai H, Lee J, Rafalski JA, Klein TM;  
XX  
DR WPI: 2000-431599/37.  
XX P-PSDB; AAY99819.  
PT Polynucleotides encoding plant protein phosphatase useful for  
XX modulating reversible protein phosphorylation in plants -  
PS Claim 4; Page 53-54; 73pp; English.

The present sequence encodes corn protein phosphatase 2A regulatory subunit A. The sequence was identified in clone p0018.chsuq10r.fis of a cDNA library made from corn ear shoot. BLAST analysis showed that the present sequence encodes protein phosphatase 2A regulatory subunit A. The sequence may be used for the recombinant production of the protein in vivo, e.g. via a gene therapy protocol, or in vitro, e.g. in fermentation culture. The protein may then be used to modulate the process of reversible protein phosphorylation in plants. It may be used directly to supplement a plant's own production of the enzyme or to rectify mutations that result in the expression of inactive protein. The protein may also be used to test for modulators of protein phosphorylation which may be used to alter the activity of the enzyme.

SQ Sequence 2061 BP; 549 A; 432 C; 498 G; 582 T; 0 other;

Query Match 100.0%; Score 121; DB 21; Length 2061;  
Best Local Similarity 100.0%; Pred. No. 2,66-33;  
Matches 121; Conservative 0; Mismatches 0; Indels 0; Gaps 0

OY 1 GTTTTCGGTCTAGCCTCGGTGACAGATCGACGCTGCCCATCTGATAATGAGCGTGCC 60  
|||  
Db 1908 GTTTCGCTCTAGCCTCGGTGACAGATCGACGCTGCCCATCTGATAATGAGCGTGCC 1967  
OY 61 TGATCCATTGTCGTTGTTTATTAATGTTGTAATTTAGCAGCACACAACGTAAGT 120  
|||||  
Db 1968 TGATCCATTGTCGTTGTTTATTAATGTTGTAATTTAGCAGCACACAACGTAAGT 2027  
OY 121 T 121  
|  
Db 2028 T 2028

RESULT 2  
ID AAA48576 standard; cDNA: 2604 BP.  
XX AAA48576;  
XX  
DT 19-SEP-2000 (first entry)  
XX  
DE cDNA encoding wheat protein phosphatase 2A regulatory subunit A.  
XX  
KW Wheat; protein phosphatase 2A; protein phosphorylation modulation;  
KM transgenic plant; gene therapy; ss.  
XX  
OS Triticum aestivum.  
XX  
FH key Location/Qualifiers  
FT CDS 101..2028  
FT /tag= a  
FT /product= "protein phosphatase 2A regulatory  
FT subunit A"  
FT /note= "contains an intron"  
FT exon 101..1537  
FT /\*tag= b  
FT /\*number= 1  
FT intron 1538..1794  
FT /\*tag= c  
FT /\*number= 1

wheat

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FT      exon          1795..2028
ET      /tag=         d
TT      /number=     2
PN      WO200036121-AZ.
PD      22-JUN-2000.
PF      15-DEC-1999;   99WO-US29823.
PR      16-DEC-1998;   98US-0112541.
PA      (DUPO ) DU PONT DE NEMOURS & CO E I.
PL      Famodu OO, Miao G, Sakai H, Lee J, Rafalski JA, Klein TM;
XX      WPI: 2000-431599/37.
DR      P-PSDB; AAV99821.
PT      Polynucleotides encoding plant protein phosphatase useful for
PS      modulating reversible protein phosphorylation in plants -
XX      Claim 4; Page 59; 73pp; English.
CC      The present sequence encodes wheat protein phosphatase 2A regulatory
CC      subunit A. The sequence was identified in clone wrl.pX0030.b10:fls of a
CC      cDNA library made from the root of a seven-day-old wheat seedling.
CC      BLAST analysis showed that the present sequence encodes protein
CC      phosphatase 2A regulatory subunit A. The sequence may be used
CC      for the recombinant production of the protein in vivo, e.g. via a gene
CC      therapy protocol, or in vitro, e.g. in fermentation culture. The protein
CC      may then be used to modulate the process of reversible protein
CC      phosphorylation in plants. It may be used directly to supplement a
CC      plant's own production of the enzyme or to rectify mutations that result
CC      in the expression of inactive protein. The protein may also be used to
CC      test for modulators of protein phosphorylation which may be used to
CC      alter the activity of the enzyme.
XX      Sequence 2604 BP; 779 A; 515 C; 579 G; 731 T; 0 other:
SQ
Query Match           24.1%; Score 29.2; DB 21; Length 2604;
Best Local Similarity 75.8%; Pred. No. 2.9;
Matches 50; Conservative 0; Mismatches 13; Indels 3; Gaps 1;
QY      38 CCATGCTGATAAATGGACGGTCCTGATCCATTGTTCGTTGTTATTAATGTTGTAAT 97
        ||| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
DB      2330 CCATGCTGATAATGATGGAGTCTGTCCCTTT---TGTTGTATTAATGTTGTAAT 2386
OY      98 TGAGCA 103
        ||| | | | | |
DB      2387 TGAGCA 2392
RESULT 3
ABQ25086
ID      ABQ25086 standard; DNA: 705 BP.
XX      ABO25086;
XX      12-JUL-2002 (first entry)
DE      Oligonucleotide for detecting cytosine methylation SEQ ID NO 11677.
KW      Human; cytosine methylation; 5'-CPG-3'; uracil; cytosine; diagnosis;
KW      drug; side effect; cancer; central nervous system; cardiovascular;
KW      gastrointestinal; respiratory system; single nucleotide polymorphism;
KW      SNP; cell differentiation; ds.
OS      Homo sapiens.
XX      WO200218632-A2.
XX      07-MAR-2002.

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XX 01-SEP-2001; 2001WO-EPI10074.
PF 01-SEP-2000; 2000DE-1043826.
XX 05-SEP-2000; 2000DE-1044543.
PR 05-SEP-2000; 2000DE-1044543.
XX (EPIC-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K, Guetig D;
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful
XX for diagnosis and prognosis, comprises selective hybridization of
XX PT amplicons from chemically treated DNA
XX
XX PS Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
XX This invention describes a novel method for determining the degree of
XX methylation of a particular cytosine in a motif 5'-CPG-3', present in a
XX genomic sample of DNA. The sample is treated chemically to convert
XX cytosine (C) but not methylated C, to uracil, then part of the genomic
XX DNA that contains the target C is amplified to form a labeled amplicon.
XX The amplicon is hybridised to two classes, each with at least one
XX member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
XX and the degree of hybridisation to both classes is determined from the
XX label on the amplicon. From the ratio of labels hybridised to the two
XX classes of oligomers, the degree of methylation is calculated. The method
XX is used: (i) for diagnosis and/or prognosis of side effects of
XX therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
XX of the central nervous, cardiovascular, gastrointestinal and respiratory
XX systems etc., particularly by detecting mutations or single nucleotide
XX polymorphisms (SNP's); and (ii) for differentiation of cell or tissue
XX types and for investigating cell differentiation. The method allows the
XX methylation status of many C residues to be determined simultaneously.
XX AB013410-AB034121 represent genomic DNA sequences used to illustrate the
XX method for determining the degree of cytosine methylation described in
XX the disclosure of the invention.
XX
XX Sequence 705 BP; 156 A; 73 C; 216 G; 260 T; 0 other;
XX
XX Query Match 23.8%; Score 28.8; DB 24; Length 705;
XX Best Local Similarity 58.0%; Pred. No. 2.5;
XX Matches 51; Conservative 0; Mismatches 37; Indels 0; Gaps 0;
XX
XX Oy 26 GATCGACCGCTGCCATGAATGAATGAGCGTCTGATCCATGTCGTTGTTATTA 85
XX ||||| | | | | | | | | | | | | | | | | | | | | | | | |
XX Db 469 GATCGAGGTGGTATATTGTAATAGATGTTGATGATGTCGTGTTGATATTATT 528
XX
XX Oy 86 ATGTTGTAATTAATTGAGCAGACACACA 113
XX || | | | | | | | | | | | | | | | | | | | | | | |
XX Db 529 TAGTATTGTTAAACGATGTGGCGCAATA 556
XX
XX RESULT 4
XX AB025087/c
XX ID AB025087 standard; DNA; 705 BP.
XX
XX AC AB025087;
XX
XX DT 12-JUL-2002 (first entry)
XX
XX DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 11678.
XX
XX KW Human; cytosine methylation; 5'-CPG-3'; uracil; cytosine; diagnosis;
XX drug; side effect; cancer; central nervous system; cardiovascular;
XX gastrointestinal; respiratory system; single nucleotide polymorphism;
XX SNP; cell differentiation; ds.
XX
XX OS Homo sapiens.
XX
XX PN WO200218632-A2.
XX

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PD 07-MAR-2002.
XX
XX 01-SEP-2001; 2001WO-EPI10074.
PF 01-SEP-2000; 2000DE-1043826.
XX 05-SEP-2000; 2000DE-1044543.
PR 05-SEP-2000; 2000DE-1044543.
XX (EPIC-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K, Guetig D;
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful
XX for diagnosis and prognosis, comprises selective hybridization of
XX PT amplicons from chemically treated DNA
XX
XX PS Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
XX This invention describes a novel method for determining the degree of
XX methylation of a particular cytosine in a motif 5'-CPG-3', present in a
XX genomic sample of DNA. The sample is treated chemically to convert
XX cytosine (C) but not methylated C, to uracil, then part of the genomic
XX DNA that contains the target C is amplified to form a labeled amplicon.
XX The amplicon is hybridised to two classes, each with at least one
XX member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers
XX and the degree of hybridisation to both classes is determined from the
XX label on the amplicon. From the ratio of labels hybridised to the two
XX classes of oligomers, the degree of methylation is calculated. The method
XX is used: (i) for diagnosis and/or prognosis of side effects of
XX therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders
XX of the central nervous, cardiovascular, gastrointestinal and respiratory
XX systems etc., particularly by detecting mutations or single nucleotide
XX polymorphisms (SNP's); and (ii) for differentiation of cell or tissue
XX types and for investigating cell differentiation. The method allows the
XX methylation status of many C residues to be determined simultaneously.
XX AB013410-AB034121 represent genomic DNA sequences used to illustrate the
XX method for determining the degree of cytosine methylation described in
XX the disclosure of the invention.
XX
XX Sequence 705 BP; 260 A; 216 C; 73 G; 156 T; 0 other;
XX
XX Query Match 23.8%; Score 28.8; DB 24; Length 705;
XX Best Local Similarity 58.0%; Pred. No. 2.5;
XX Matches 51; Conservative 0; Mismatches 37; Indels 0; Gaps 0;
XX
XX Oy 26 GATCGACCGCTGCCATGAATGAATGAGCGTCTGATCCATGTCGTTGTTATTA 85
XX ||||| | | | | | | | | | | | | | | | | | | | | | | | |
XX Db 237 GATCGAGGTGGTATATTGTAATAGATGTTGATGATGTCGTGTTGATATTATT 178
XX
XX Oy 86 ATGTTGTAATTAATTGAGCAGACACACA 113
XX || | | | | | | | | | | | | | | | | | | | | | | |
XX Db 177 TAGTATTGTTAAACGATGTGGCGCAATA 150
XX
XX RESULT 5
XX AAV64373
XX ID AAV64373 standard; cDNA; 1519 BP.
XX
XX AC AAV64373;
XX
XX DT 15-FEB-1999 (first entry)
XX
XX DE GABA-gated chloride channel TBW-a3 cDNA.
XX
XX KW TBW-a3; GABA-gated chloride channel; tobacco budworm; insecticide;
XX ss.
XX
XX OS Heliothis virescens.
XX
XX PH Key
XX FT CDS 1..1446
XX /*tag= a
XX

```

FT sig\_peptide 1..66  
 FT /\*tag= b  
 FT sig\_peptide 67..143  
 FT /\*tag= c  
 XX  
 PN WO9849185-A1.  
 XX  
 PD 05-NOV-1998.  
 XX  
 XX 27-APR-1998; 98WO-US08563.  
 PF  
 XX 02-JAN-1998; 98US-0002361.  
 PR 28-APR-1997; 97US-0044976.  
 XX  
 PA (FMCC ) FMC CORP.  
 XX  
 PI Halling BP, Yuhas DA;  
 XX  
 DR WPI: 1999-009411/01.  
 XX P-PSDB; AAW81635-36.  
 XX  
 PT New isolated Lepidoptera GABA-gated chloride channels - comprise 3  
 PT isoforms isolated from the tobacco budworm *Heliothis virescens*, used  
 PT for characterizing bioactive agents, e.g. insecticides  
 PS  
 PS Claim 1d; Fig 2; 55pp; English.  
 XX  
 CC This cDNA sequence encompasses the open reading frame encoding  
 CC GABA-gated chloride channel TBW-a3 (see AAW81635-36) of tobacco  
 CC budworm (*Heliothis virescens*). TBW-a3, TBW-a2 (see AAW81633-34)  
 CC and TBW-a1 (see AAW81637) proteins are 3 receptor isoforms that show  
 CC sequence homology to each other and to other insect GABA-gated  
 CC chloride channels. TBW-2a cDNA was obtained from *H. virescens* 4th  
 CC instar larva RNA by PCR and RACE amplifications. The invention  
 CC provides expression vectors in which a nucleic acid encoding a  
 CC GABA-gated chloride channel is driven by an inducible promoter, and  
 CC a claimed process for producing a GABA-gated chloride channel by  
 CC transformed cells. The GABA-gated channels or cells expressing  
 CC them can be used for characterizing a bioactive agent (claimed),  
 CC e.g. for use as an insecticide. Probes and primers that identify  
 CC or amplify GABA-gated chloride channel nucleic acids of the  
 CC invention are also claimed.  
 CC  
 CC  
 SQ Sequence 1519 BP; 421 A; 362 C; 347 G; 388 T; 1 other:  
 Query Match 23.8%; Score 28.8; DB 20; Length 1519;  
 Best Local Similarity 52.5%; Pred. No. 3.3;  
 Matches 63; Conservative 0; Mismatches 57; Indels 0; Gaps 0;  
 OY 2 TTTTCGCTACCTCGGTGGACAGATCGACGCTGCCATGCTGATAATGACGGTCT 61  
 DB 253 TTTTACTTACAGACATTTTGGACTGATCCTCGATTAGCAAAAAGAACCGAGTT 312  
 OY 62 GATCCATTGCTGCTGTTTATTAATGTTATGAGACAGACACAGTACGTT 121  
 DB 313 GAAACTTATCTGTGGCTCAGATTCAATAAGAACATATGAGTACCGACGCTTCTT 372

PN WO200160860-A2.  
 XX  
 PD 23-AUG-2001.  
 XX  
 PF 20-FEB-2001; 2001WO-US05171.  
 XX  
 PR 17-FEB-2000; 2000US-183319P.  
 PR 16-MAR-2000; 2000US-189862P.  
 PR 25-MAY-2000; 2000US-207454P.  
 PR 09-JUN-2000; 2000US-211314P.  
 PR 18-JUL-2000; 2000US-219007P.  
 PR 13-DEC-2000; 2000US-235281P.  
 XX  
 PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
 XX  
 PI Schlegel R, Endege WO, Monahan JE;  
 XX  
 DR WPI: 2001-662795/76.  
 XX  
 PT Novel isolated nucleic acid molecule associated with cancerous state of  
 PT prostate cells and correlating with presence of prostate cancer, useful  
 PT for detecting presence of prostate cancer, stage of prostate cancer -  
 PS  
 PS Claim 1; Page 9499; 11750pp; English.  
 XX  
 CC The invention relates to an isolated nucleic acid molecule (I) comprising  
 CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
 CC specification or its complement. (I) is useful for:  
 CC (a) assessing whether a patient is afflicted with prostate cancer;  
 CC (b) monitoring the progression of prostate cancer in a patient;  
 CC (c) assessing the efficacy of a test compound to inhibit prostate  
 CC cancer in a patient;  
 CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer  
 CC in a patient;  
 CC (e) selecting a composition for inhibiting prostate cancer in a patient;  
 CC (f) assessing the prostate cell carcinogenic potential of a compound;  
 CC (g) determining whether prostate cancer has metastasized in a patient;  
 CC (h) assessing the aggressiveness or indolence of prostate cancer in a  
 CC patient;  
 CC (I) is also useful as a pharmacodynamic or pharmacogenomic marker.  
 CC  
 CC  
 SQ Sequence 405 BP; 106 A; 86 C; 102 G; 111 T; 0 other:  
 Query Match 23.6%; Score 28.6; DB 23; Length 405;  
 Best Local Similarity 59.0%; Pred. No. 2.4;  
 Matches 49; Conservative 0; Mismatches 34; Indels 0; Gaps 0;  
 OY 9 TCTACCCGCTGGACACATGACGCTGCCATGCTGATAATGACGGTCTGATCAT 68  
 DB 321 TATAGCGACAGTGCATGTCACCTTGATGATGCTGAAGAAAAACCGTACCATCAT 380  
 OY 69 TGTTCGTGTGTATTAATGTTG 91  
 DB 381 TCTTCATGCTGTGGTGGAGTTG 403

RESULT 6  
 ABV48454  
 ID ABV48454 standard; cDNA; 405 BP.  
 XX  
 AC ABV48454;  
 XX  
 DT 17-SEP-2002 (first entry)  
 XX  
 DE Human prostate expression marker cDNA 48445.  
 XX  
 KW Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
 KW pharmacogenomic marker; gene; ss.  
 XX  
 OS Homo sapiens.  
 XX

RESULT 7  
 AA088698  
 ID AA088698 standard; cDNA to mRNA; 297 BP.  
 XX  
 AC AA088698;  
 XX  
 DT 11-NOV-1995 (first entry)  
 XX  
 DE Beta-amyloid precursor protein C-terminal peptide mutant gene.  
 XX  
 KW Human; beta-amyloid precursor protein mutant; C-terminal peptide;  
 KW gene transfer; transgenic animal; Alzheimer disease model;  
 KW gene therapy; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN EP653154-A.

XX 17-MAY-1995.  
 PD 07-NOV-1994; 94EP-0117512.  
 XX 12-NOV-1993; 93JP-0306026.  
 PR (FARH ) HOECHST JAPAN LTD.  
 PA (FARH ) HOECHST JAPAN KK.  
 XX Kawayabayashi T, Kobayashi T, Sato M, Shoji M, Tada N;  
 PI WPI: 1995-180492/24.  
 DR P-PSDB: AAR/4696.  
 XX  
 PT Transgenic animal model for Alzheimer's disease - contains DNA encoding  
 PT part of beta-amyloid precursor protein in a gene construct designed for  
 PT over-expression in various cell types  
 PS  
 PS Claim 2; Page 13-14; 32pp; English.  
 XX  
 CC The sequence encodes a human brain beta-amyloid precursor protein  
 CC (APP) mutant C-terminal peptide, and differs from AA088696 by  
 CC conversion of Val to Ile at codon 46. The DNA may be transferred  
 CC along with an APP signal peptide gene (e.g. AA088695) into somatic and  
 CC germ cells of a non-human mammal, and the resulting transgenic animal  
 CC may be used as a model for Alzheimer disease (AD). The animal model  
 CC exhibits symptoms similar to AD, producing large quantities of APP  
 CC C-terminal peptide, death of neuron cells in pyramidal cells at  
 CC cerebral amyloid regions, increases in glial cells and deposition  
 CC of abnormally phosphorylated tau protein. The animal model may  
 CC be used to develop new therapies for AD, including gene therapy  
 CC strategies.  
 CC  
 SO Sequence 297 BP; 87 A; 64 C; 77 G; 69 T; 0 other;  
 XX  
 Query Match 23.3%; Score 28.2; DB 16; Length 297;  
 Best Local Similarity 59.3%; Pred. No. 3.1;  
 Matches 48; Conservative 0; Mismatches 33; Indels 0; Gaps 0;  
 QY 11 TAGCCTGGTGGACAGATCGACGCTGATTAATGAGACGGTCTGATCCATTG 70  
 Db 122 TAGGACAGTATCATCATCTGCTGGTGTGATGCTGAGAGAAACAGTACATCCATTG 181  
 QY 71 TTGCTGTGTTATTATGTTG 91  
 Db 182 ATCATGTGTGCTGTGAGGTTG 202  
 XX  
 RESULT 8  
 ID AAC9387 standard; DNA; 300 BP.  
 AC AAC9387;  
 XX  
 DT 27-JUN-2002 (first entry)  
 XX  
 DE DNA of wild type C99 portion of human APP with the London mutation.  
 XX  
 KW Neuroprotective; nootropic; transgenic fly; Alzheimer's disease; Abeta;  
 KW amyloid precursor protein; tissue-specific expression control; human APP;  
 KW APP pathway modulator; gene therapy; mutant; ds.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200226820-A2.  
 PD 04-APR-2002.  
 XX  
 PF 01-OCT-2001; 2001WO-EP11345.  
 XX  
 PR 29-SEP-2000; 2000US-236893P.  
 PR 14-JUN-2001; 2001US-298309P.  
 XX

XX (NOVS ) NOVARTIS AG.  
 PA (NOVS ) NOVARTIS-ERFINDUNGEN VERN GES MBH.  
 XX  
 PI Cohen D, Dengler UJ, Finelli AL, Freuler F, Konsolaki M;  
 PI Reinhardt MWM, Zisman S;  
 XX WPI: 2002-315796/35.  
 DR  
 XX  
 PT New transgenic fly, containing DNA encoding an Abeta portion of human  
 PT APP, useful for identifying agents which modulate the APP pathway and  
 PT which can be used to treat Alzheimer's disease -  
 XX  
 PS Claim 4; Page 84; 129pp; English.  
 XX  
 CC The invention relates to a transgenic fly whose genome comprises DNA  
 CC encoding a polypeptide having the Abeta portion of human amyloid  
 CC precursor protein (APP), fused to a signal sequence. The DNA sequence  
 CC encodes a 123 (Abeta40) or 129 (Abeta42) amino acid sequence, given in  
 CC the specification. The DNA sequence is operably linked to a tissue-  
 CC specific expression control sequence. Expression of the sequence gives  
 CC the fly an altered phenotype. The purpose of the invention is for  
 CC identifying agents that inhibit or promote the expression and/or function  
 CC of genes or encoded polypeptides which modify the APP pathway. The agent  
 CC is a compound, triple helix DNA, antisense oligonucleotide, double  
 CC stranded RNA molecule, ribozyme, or particularly an antibody. It is used  
 CC to treat conditions such as Alzheimer's disease. The agent can be used as  
 CC an APP pathway modulator or in gene therapy. This polynucleotide sequence  
 CC represents the DNA of the wild type C99 portion of human APP with the  
 CC London mutation of the invention.  
 CC  
 SO Sequence 300 BP; 88 A; 65 C; 78 G; 69 T; 0 other;  
 XX  
 Query Match 23.3%; Score 28.2; DB 24; Length 300;  
 Best Local Similarity 59.3%; Pred. No. 3.1;  
 Matches 48; Conservative 0; Mismatches 33; Indels 0; Gaps 0;  
 QY 11 TAGCCTGGTGGACAGATCGACGCTGATTAATGAGACGGTCTGATCCATTG 70  
 Db 122 TAGGACAGTATCATCATCTGCTGGTGTGATGCTGAGAGAAACAGTACATCCATTG 181  
 QY 71 TTGCTGTGTTATTATGTTG 91  
 Db 182 ATCATGTGTGCTGTGAGGTTG 202  
 XX  
 RESULT 9  
 ID AAC79021 standard; DNA; 1357 BP.  
 AC AAC79021;  
 XX  
 DT 14-FEB-2001 (first entry)  
 XX  
 DE Human secreted protein gene 25 clone HODAH24.  
 XX  
 KW Cytostatic; immunosuppressive; nootropic; neuroprotective; antiviral;  
 KW antiallergic; hepatoprotective; antidiabetic; antiinflammatory; antitumor;  
 KW vulnerary; anticonvulsant; antibacterial; antifungal; antiparasitic;  
 KW cardiant; gene therapy; cancer; immune disorder; cardiovascular disorder;  
 KW neurological disease; infection; human; secreted protein; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200058358-A1.  
 PD 05-OCT-2000.  
 XX  
 PF 23-MAR-2000; 2000WO-US07725.  
 XX  
 PR 26-MAR-1999; 99US-0126602.  
 PR 14-JAN-2000; 2000US-0176063.  
 XX

PA (HUMA-) HUMAN GENOME SCI INC.  
 XX  
 PI Rosen CA, Ruben SM, Komatsoulis G;  
 XX  
 DR WPI: 2000-594640/56.  
 DR P-PSDB: AAB44359.  
 XX  
 PT Fourty nine nucleic acid molecules encoding human secreted proteins,  
 PT useful in the prevention, treatment and diagnosis of cancer, immune  
 PT disorders, cardiovascular disorders and neurological diseases -  
 XX  
 PS Claim 1: Page 327-328; 367pp; English.  
 XX  
 CC The invention relates to the isolation of genes AAC78997-C79045 encoding  
 CC 49 human secreted proteins AAB4435-844382. The genes can be used to  
 CC generate fusion proteins by linking to the gene for the human  
 CC immunoglobulin G Fc portion (AAC78988) for increasing the stability of  
 CC the fusion protein as compared to the human protein only. The genes and  
 CC proteins are useful for preventing, ameliorating or treating medical  
 CC conditions, e.g. by protein or gene therapy. The genes are isolated  
 CC from a range of human tissues disclosed in the specification. The  
 CC nucleic acids, proteins, antibodies and (ant)agonists are useful in  
 CC the diagnosis, treatment and prevention of: (a) cancer, e.g. breast  
 CC and ovarian cancer, and other cancers of the adrenal gland, bone, bone  
 CC marrow, breast, gastrointestinal tract, liver, lung, or urogenital;  
 CC (b) immune disorders e.g. Addison's disease, allergies, autoimmune  
 CC haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's  
 CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative  
 CC colitis; (c) cardiovascular disorders such as myocardial ischaemias; (d)  
 CC wound healing; (e) neurological diseases e.g. cerebral anoxia and  
 CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal  
 CC and parasitic infections.  
 XX  
 SQ Sequence 1357 BP; 386 A; 272 C; 299 G; 398 T; 2 other:  
 XX  
 Query Match 23.1%; Score 28; DB 21; Length 1357;  
 Best Local Similarity 52.6%; Pred. No. 6.1;  
 Matches 61; Conservative 0; Mismatches 55; Indels 0; Gaps 0;  
 OY 1 GTTTTCGCTAGCCTGGTGGACAGATGACGCTGCCATGCTGTAATGAGCGTCC 60  
 DB 871 GTTACCTTTGTGACGTGATCCACCCACCTGGCTCCCAAGTGCTGAGATTACAGAGC 930  
 OY 61 TGATCCATGCTGCTGTGTATATATGTTGATATAGCAGACACACACGT 116  
 DB 931 TGAGCCATGTCGCCCGCATTTTAAATTTTAAATTAATTGACACACCT 986  
 RESULT 10  
 ABL04431/C  
 ID ABL04431 standard; cDNA; 1086 BP.  
 XX  
 AC ABL04431;  
 XX  
 DT 26-MAR-2002 (first entry)  
 XX  
 DE Drosophila melanogaster expressed polynucleotide SEQ ID NO 7775.  
 XX  
 KW Drosophila; developmental biology; cell signalling; insecticide;  
 KW pharmaceutical; gene; ss.  
 XX  
 OS Drosophila melanogaster.  
 XX  
 PN WO200171042-A2.  
 XX  
 PD 27-SEP-2001.  
 XX  
 PF 23-MAR-2001; 2001WO-US09231.  
 XX  
 PR 23-MAR-2000; 2000US-191637P.  
 PR 11-JUL-2000; 2000US-0614150.  
 XX  
 PA (PEKE ) PE CORP NY.

XX  
 PI Venter JC, Adams M, Li PWD, Myers EW;  
 XX  
 DR WPI: 2001-656860/75.  
 DR P-PSDB: ABB60328.  
 XX  
 PT New isolated nucleic acid detection reagent for detecting 1000 or more  
 PT genes from Drosophila and for elucidating cell signalling and cell-cell  
 PT interactions -  
 XX  
 PS Claim 1: SEQ ID NO 7775; 21pp + Sequence Listing; English.  
 XX  
 CC The invention relates to an isolated nucleic acid detection reagent  
 CC capable of detecting 1000 or more genes from Drosophila. The invention is  
 CC useful in developmental biology and in elucidating cell signalling and  
 CC cell-cell interactions in higher eukaryotes for the development of  
 CC insecticides, therapeutics and pharmaceutical drugs. The invention  
 CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA  
 CC sequences (ABB57737-ABB72072).  
 CC (ABB57737-ABB72072).  
 CC The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pct\_sequences.  
 XX  
 SQ Sequence 1086 BP; 295 A; 267 C; 273 G; 251 T; 0 other:  
 XX  
 Query Match 22.8%; Score 27.6; DB 23; Length 1086;  
 Best Local Similarity 60.8%; Pred. No. 7.9;  
 Matches 45; Conservative 0; Mismatches 29; Indels 0; Gaps 0;  
 OY 36 GCCCATGCTGAATAGACGCTCTGATCCATGTTGCTGTTATGTTGATATA 95  
 DB 943 GCCCATATCCGTAACGTGAGGTCCTCCGCTAGACGTTATCCGCTAGTAATATAA 884  
 OY 96 ATGAGCAGACAC 109  
 DB 883 ATGATGACGACCC 870  
 RESULT 11  
 ABL04430  
 ID ABL04430 standard; cDNA; 3205 BP.  
 XX  
 AC ABL04430;  
 XX  
 DT 26-MAR-2002 (first entry)  
 XX  
 DE Drosophila melanogaster expressed polynucleotide SEQ ID NO 7772.  
 XX  
 KW Drosophila; developmental biology; cell signalling; insecticide;  
 KW pharmaceutical; gene; ss.  
 XX  
 OS Drosophila melanogaster.  
 XX  
 PN WO200171042-A2.  
 XX  
 PD 27-SEP-2001.  
 XX  
 PF 23-MAR-2001; 2001WO-US09231.  
 XX  
 PR 23-MAR-2000; 2000US-191637P.  
 PR 11-JUL-2000; 2000US-0614150.  
 XX  
 PA (PEKE ) PE CORP NY.  
 XX  
 PI Venter JC, Adams M, Li PWD, Myers EW;  
 XX  
 DR WPI: 2001-656860/75.  
 DR P-PSDB: ABB60327.  
 XX  
 PT New isolated nucleic acid detection reagent for detecting 1000 or more  
 PT genes from Drosophila and for elucidating cell signalling and cell-cell  
 PT interactions -



xx Claim 1; SEQ ID NO 7772; 21pp + Sequence Listing; English.  
xx  
xx  
xx The invention relates to an isolated nucleic acid detection reagent  
cc capable of detecting 1000 or more genes from Drosophila. The invention is  
cc useful in developmental biology and in elucidating cell signalling and  
cc cell-cell interactions in higher eukaryotes for the development of  
cc insecticides, therapeutics and pharmaceutical drugs. The invention  
cc discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA  
cc sequences (ABL01840-ABL16175) and the encoded proteins  
cc (ABB57737-ABB72072).  
cc The sequence data for this patent did not form part of the printed  
cc specification, but was obtained in electronic format directly from WIPO  
cc at ftp.wipo.int/pub/published\_pct\_sequences.  
xx  
xx Sequence 3205 BP; 932 A; 681 C; 655 G; 937 T; 0 other;  
SQ  
Query Match 22.8%; Score 27.6; DB 23; Length 3205;  
Best Local Similarity 60.8%; Pred. No. 12;  
Matches 45; Conservative 0; Mismatches 29; Indels 0; Gaps 0;  
OY 36 GCCCATGCTGATTAATGACGCTCCTGATTCATTTCTGTGTATTATTAATGTTGATA 95  
DB 1199 GCCCATATCCGTAACCTGAGAGGCTCCCTCCGTAGACGTTATTCGGCTAGTATATAA 1258  
OY 96 ATTGACGACGACAC 109  
DB 1259 ATTGATCAGCACCC 1272  
RESULT 12  
ABL04426  
ID ABL04426 standard; cDNA; 3457 BP.  
xx  
xx ABL04426;  
xx  
xx 26-MAR-2002 (first entry)  
xx  
xx Drosophila melanogaster expressed polynucleotide SEQ ID NO. 7760.  
xx  
xx Drosophila; developmental biology; cell signalling; insecticide;  
xx pharmaceutical; gene; ss.  
xx Drosophila melanogaster.  
xx  
xx WO200171042-A2.  
xx  
xx 27-SEP-2001.  
xx  
xx 23-MAR-2001; 2001WO-US09231.  
xx  
xx 23-MAR-2000; 2000US-191637P.  
xx 11-JUL-2000; 2000US-0614150.  
xx  
xx (PEKE ) PE CORP NY.  
xx  
xx Venter JC, Adams M, LI PWD, Myers EW;  
xx  
xx WPI: 2001-656860/75.  
xx P-PSDB: ABB60323.  
xx  
xx New isolated nucleic acid detection reagent for detecting 1000 or more  
xx genes from Drosophila and for elucidating cell signalling and cell-cell  
xx interactions -  
xx  
xx Claim 1; SEQ ID NO 7760; 21pp + Sequence Listing; English.  
xx  
xx The invention relates to an isolated nucleic acid detection reagent  
cc capable of detecting 1000 or more genes from Drosophila. The invention is  
cc useful in developmental biology and in elucidating cell signalling and  
cc cell-cell interactions in higher eukaryotes for the development of  
cc insecticides, therapeutics and pharmaceutical drugs. The invention  
cc discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA  
cc sequences (ABL01840-ABL16175) and the encoded proteins  
cc (ABB57737-ABB72072).  
cc The sequence data for this patent did not form part of the printed  
cc specification, but was obtained in electronic format directly from WIPO  
cc at ftp.wipo.int/pub/published\_pct\_sequences.  
xx  
xx Sequence 4659 BP; 1472 A; 1196 C; 1153 G; 838 T; 0 other;  
SQ  
Query Match 22.6%; Score 27.4; DB 23; Length 4659;  
Best Local Similarity 69.8%; Pred. No. 16;

cc sequences (ABL01840-ABL16175) and the encoded proteins  
cc (ABB57737-ABB72072).  
cc The sequence data for this patent did not form part of the printed  
cc specification, but was obtained in electronic format directly from WIPO  
cc at ftp.wipo.int/pub/published\_pct\_sequences.  
xx  
xx Sequence 3457 BP; 895 A; 779 C; 690 G; 1093 T; 0 other;  
SQ  
Query Match 22.8%; Score 27.6; DB 23; Length 3457;  
Best Local Similarity 60.8%; Pred. No. 12;  
Matches 45; Conservative 0; Mismatches 29; Indels 0; Gaps 0;  
OY 36 GCCCATGCTGATTAATGACGCTCCTGATTCATTTCTGTGTATTATTAATGTTGATA 95  
DB 3299 GCCCATATCCGTAACCTGAGAGGCTCCCTCCGTAGACGTTATTCGGCTAGTATATAA 3358  
OY 96 ATTGACGACGACAC 109  
DB 3359 ATTGATCAGCACCC 3372  
RESULT 13  
ABL29895/C  
ID ABL29895 standard; DNA; 4659 BP.  
xx  
xx ABL29895;  
xx  
xx 26-MAR-2002 (first entry)  
xx  
xx Drosophila melanogaster genomic polynucleotide SEQ ID NO 41158.  
xx  
xx Drosophila; developmental biology; cell signalling; insecticide;  
xx pharmaceutical; gene; ds.  
xx Drosophila melanogaster.  
xx  
xx WO200171042-A2.  
xx  
xx 27-SEP-2001.  
xx  
xx 23-MAR-2001; 2001WO-US09231.  
xx  
xx 23-MAR-2000; 2000US-191637P.  
xx 11-JUL-2000; 2000US-0614150.  
xx  
xx (PEKE ) PE CORP NY.  
xx  
xx Venter JC, Adams M, LI PWD, Myers EW;  
xx  
xx WPI: 2001-656860/75.  
xx  
xx New isolated nucleic acid detection reagent for detecting 1000 or more  
xx genes from Drosophila and for elucidating cell signalling and cell-cell  
xx interactions -  
xx  
xx Claim 1; SEQ ID NO 41158; 21pp + Sequence Listing; English.  
xx  
xx The invention relates to an isolated nucleic acid detection reagent  
cc capable of detecting 1000 or more genes from Drosophila. The invention is  
cc useful in developmental biology and in elucidating cell signalling and  
cc cell-cell interactions in higher eukaryotes for the development of  
cc insecticides, therapeutics and pharmaceutical drugs. The invention  
cc discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA  
cc sequences (ABL01840-ABL16175) and the encoded proteins  
cc (ABB57737-ABB72072).  
cc The sequence data for this patent did not form part of the printed  
cc specification, but was obtained in electronic format directly from WIPO  
cc at ftp.wipo.int/pub/published\_pct\_sequences.  
xx  
xx Sequence 4659 BP; 1472 A; 1196 C; 1153 G; 838 T; 0 other;  
SQ  
Query Match 22.6%; Score 27.4; DB 23; Length 4659;  
Best Local Similarity 69.8%; Pred. No. 16;

[illegible]

XX	Human CARD-4 DNA.	
DE		
XX		
KW	CARD-3; caspase recruitment domain; CARD-4; regulation; detection;	
KW	caspase activation; detection; screening; therapy; diagnosis; disease;	
KW	apoptotic cell death; Fas/Apo-1 receptor complex; TNF receptor complex;	
KW	cancer; follicular lymphoma; carcinoma; p53 mutation; viral infection;	
KW	hormone-dependent tumor; autoimmune disorder; Alzheimer's disease;	
KW	systemic lupus erythematosus; immune-mediated glomerulonephritis; stroke;	
KW	Parkinson's disease; amyotrophic lateral sclerosis; retinitis pigmentosa;	
KW	spinal muscular dystrophy; cerebellar degeneration; anaemia; drug;	
KW	myelodysplastic syndrome; myocardial infarction; cell proliferation;	
KW	cell differentiation; cell survival; CARD-4L; CARD-4S; CARD-4Y;	
XX	CARD-4Z; human; ds.	
XX		
OS	Homo sapiens.	
XX		
PH	Key	Location/Qualifiers
FT	exon	364..685
FT		/*tag= a
FT		/number= 1
FT		485..31768
FT	CDS	/*tag= b
FT		/product= "CARD-4"
FT		686..2094
FT	intron	/*tag= c
FT		/number= 1
FT		2095..2269
FT	exon	/*tag= d
FT		/number= 2
FT		2270..4365
FT	intron	/*tag= e
FT		/product= 2
FT		4366..6190
FT	exon	/*tag= f
FT		/number= 3
FT		6191..9024
FT	intron	/*tag= g
FT		/number= 3
FT		9025..9108
FT	exon	/*tag= h
FT		/product= 4
FT		9109..10355
FT	intron	/*tag= i
FT		/number= 4
FT		10356..10439
FT	exon	/*tag= j
FT		/number= 5
FT		10440..11181
FT	intron	/*tag= k
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FT		11182..11265
FT	exon	/*tag= l
FT		/number= 6
FT		11266..19749
FT	intron	/*tag= m
FT		/number= 6
FT		19750..19833
FT	exon	/*tag= n
FT		/number= 7
FT		19834..21324
FT	intron	/*tag= o
FT		/number= 7
FT		21325..21408
FT	exon	/*tag= p
FT		/number= 8
FT		21409..24226
FT	intron	/*tag= q
FT		/number= 8
FT		24227..24310
FT	exon	/*tag= r
FT		/number= 9
FT		24311..27948
FT	intron	

FT	/*tag= s	
FT	/number= 9	
FT	27949..28032	
FT	/tag= t	
FT	/product= 10	
FT	28033..31695	
FT	/tag= u	
FT	/number= 10	
FT	31696..32024	
FT	/tag= v	
FT	/number= 11	
XX		
PN	WO9940102-A1.	
XX		
PD	12-AUG-1999:	
XX		
PF	05-FEB-1999:	99WO-US02544.
PR	08-DEC-1998:	98US-0207359.
PR	06-FEB-1998:	98US-0019942.
PR	17-JUN-1998:	98US-0099041.
XX		
PA	(MILL-) MILLENNIUM PHARM INC.	
XX		
P1	Bertin J:	
XX		
DR	WPI: 1999-494269/41.	
XX		
PT	Novel CARD-3 and CARD-4 genes and polypeptides used or treating	
PT	regulation of cellular proliferation and differentiation and cell	
XX	survival	
PS		
XX	Example 13: Fig 18; 181pp; English.	
CC	This invention describes the isolation of novel human caspase	
CC	recruitment domain, CARD-3 and CARD-4 polynucleotides and proteins and a	
CC	partial murine CARD-4L protein and genes. The genes and proteins of	
CC	the invention are involved in the regulation of caspase activation.	
CC	The caspase recruitment domain (CARD) polynucleotides, polypeptides,	
CC	homologues and antibodies can be used in screening assays, detection	
CC	assays, predictive medicine and therapeutic and prophylactic methods of	
CC	treatment. The methods may be used to diagnose and treat patients which	
CC	are suffering from a disorder associated with abnormal level or rate of	
CC	apoptotic cell death, abnormal activity of the Fas/Apo-1 receptor	
CC	complex, abnormal activity of the TNF receptor complex, or abnormal	
CC	activity of a caspase. Diseases that may be treated include cancer	
CC	(particularly follicular lymphoma, carcinomas associated with mutations	
CC	in p53 and hormone-dependent tumours), autoimmune disorders (e.g.	
CC	systemic lupus erythematosus, immune-mediated glomerulonephritis), viral	
CC	infections, Alzheimer's disease, Parkinson's disease, amyotrophic lateral	
CC	sclerosis, retinitis pigmentosa, spinal muscular dystrophy, cerebellar	
CC	degeneration, anaemia, myelodysplastic syndrome, myocardial infarction,	
CC	and stroke. CARD-3 protein interacts with other cellular proteins, and so	
CC	can be used for regulation of cellular proliferation and differentiation	
CC	and cell survival. The CARD proteins may also be used to for screen drugs	
CC	or compounds which modulate their activity. The CARD-4 gene can express a	
CC	long transcript that encodes CARD-4L, a short transcript that encodes	
CC	CARD-4S or two CARD-4 splice variants, CARD-4Y and CARD-4Z. This sequence	
CC	represents a genomic DNA sequence which encodes the human CARD-4 protein.	
XX		
SQ	Sequence 32042 BP: 7389 A: 7540 C: 7721 G: 9392 T: 0 other:	
	Query Match	22.6%: Score 27.4: DB 20: Length 32042:
	Best Local Similarity	62.3%: Pred. No. 30:
	Matches	43: Conservative 0: Mismatches 26: Indels 0: Gaps 0
QY	28 TCGACGTGCCCATCTGATTAATGACGCGTCCATGATTCGTTGTTATTAAAT 87	
Db	18335 TTGACAGTACCCATCCTATATGAGTGAGGTCCTGCTCATTTGTGTTGATTCTTGAG 18394	
QY	88 GTTGATATA 96	
Db	18395 GCTTTTATA 18403	

ID	AAAF30011 standard; cDNA; 32042 BP.
XX	AAAF30011;
DT	23-APR-2001 (first entry)
DE	Human CARD-4 gene.
XX	
KW	CARD-4; caspase recruitment domain; human; cancer; infection;
KW	autoimmune disease; neurological disease; haematological disease;
KW	immune disease; inflammation; antitumour; antiseptic;
KW	immunomodulator; antiinflammatory; apoptosis; diagnosis;
KW	gene therapy; chromosome 7; ds.
XX	
OS	Homo sapiens.
XX	
FH	Location/Qualifiers
FT	CDS
FT	485..31768
FT	/tag= a
FT	/note= "contains introns"
FT	354..685
FT	/tag= b
FT	/number= 1
FT	686..2094
FT	/tag= c
FT	/number= 1
FT	2095..2269
FT	/tag= d
FT	/number= 2
FT	2270..4365
FT	/tag= e
FT	/number= 2
FT	4366..6190
FT	/tag= f
FT	/number= 3
FT	6191..9024
FT	/tag= g
FT	/number= 3
FT	9025..9108
FT	/tag= h
FT	/number= 4
FT	9109..10355
FT	/tag= i
FT	/number= 4
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FT	/number= 5
FT	11182..11265
FT	/tag= l
FT	/number= 6
FT	11266..19749
FT	/tag= m
FT	/number= 6
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FT	/tag= n
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FT	/tag= o
FT	/number= 7
FT	21325..21408
FT	/tag= p
FT	/number= 8
FT	21409..24226
FT	/tag= q
FT	/number= 8
FT	24227..24310
FT	exon

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FT      /tag= r
FT      /number= 9
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FT      /tag= s
FT      /number= 9
FT      27949..28032
FT      /tag= t
FT      /number= 10
FT      28033..31695
FT      /tag= u
FT      /number= 10
FT      31696..32024
FT      /tag= v
FT      /number= 11
XX      WO200100826-A2.
XX      04-JAN-2001.
XX      PD
XX      28-JUN-2000; 2000MO-US17691.
XX      PR
XX      28-JUN-1999; 99US-0340620.
XX      PA
XX      (MILL-) MILLENNIUM PHARM INC.
XX      PI
XX      Bertin J;
XX      WPI; 2001-061973/07.
XX      DR      P-PSDB; AAB20080, AAB20081, AAB20082, AAB20083.
XX      PT      Isolated intracellular proteins predicted to be involved in regulating
XX      PT      caspase activation are used for diagnosis and treatment of e.g. cancer,
XX      PT      viral infections, autoimmune diseases, neurological diseases and
XX      PT      haematological disorders -
XX      PS
XX      Example 13; Fig 18; 208pp; English.
XX
CC      The present sequence is that of the human caspase recruitment
CC      domain 4 (CARD-4) gene on chromosome 7. CARD-4 exists in at least
CC      CC      4 forms, i.e. the long form CARD-4L (see AAB20080), the short form
CC      CARD-4S (see AAB20081), and splice variants CARD-4Y (see AAB20082)
CC      and CARD-4Z (see AAB20082). It is an intracellular protein
CC      CC      predicted to be involved in regulating caspase activation. It
CC      activates the NF-kappaB pathway and enhances caspase 9-mediated
CC      cell death. Methods of diagnosing and treating patients suffering
CC      from a disorder associated with an abnormal level or rate of apoptotic
CC      cell death, abnormal activity of the Fas/Apo-1 receptor complex,
CC      abnormal activity of the tumour necrosis factor receptor complex,
CC      or abnormal activity of a caspase involve administering a compound
CC      that modulates the expression or activity of CARD-3, CARD-4, CARD-5
CC      or CARD-6 e.g. a small molecule, antisense nucleic acid, ribozyme
CC      or polypeptide. Such disorders include cancer, viral infection,
CC      autoimmune disorders, neurological diseases, haematological
CC      disorders, inflammatory disorders and immune disorders. The CARD-4
CC      gene is useful for genetic information and mapping and identifying
CC      mutations, e.g. mutations in splice donor or acceptor sites.
XX      XX
XX      Sequence 32042 BP; 7389 A; 7540 C; 7721 G; 9392 T; 0 other;
SO

```

Query Match 22.6%; Score 27.4; DB 22; Length 32042;  
 Best Local Similarity 62.3%; Pred. No. 30;  
 Matches 43; Mismatches 0; Indels 0; Gaps 0;

```

OY      28 TCGAGCGTCCCATGCTAATAATGACGCGCTCATGCTGCTGTTATTAAT 87
DB      18335 TTGACAGTACCCATCTCAATGAGTGTGAGAGTCTCTGCTATTTGCTTGG 18394
OY      88 GTTGATATA 96
DB      18395 GCTTTTAA 18403

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AAL40765
ID      AAL40765 standard; DNA; 32042 BP.
XX
AC      AAL40765;
XX
DT      27-SEP-2002 (first entry)
XX
DE      Genomic DNA of human CARD-4 SEQ ID No 44.
XX
XX      Cytostatic; virucide; immunomodulatory; neuroprotective; antialzheimers;
XX      neurotrophic; antianemic; caspase recruitment domain; CARD; CARD-4L; p3;
XX      cancer; CARD-4S; follicular lymphoma; carcinoma; autoimmune disorder;
XX      hormone-dependent tumour; breast cancer; prostate cancer; ovarian cancer;
XX      systemic lupus; herpes virus; poxvirus; adenovirus; neurological disease;
XX      anaemia; neutropenia; myelodysplastic syndrome; human; ds.
XX
XX      Homo sapiens.
XX
XX      US6369196-B1.
XX      PD
XX      09-APR-2002.
XX
XX      05-FEB-1999; 99US-0245281.
XX      PF
XX      06-FEB-1998; 98US-0019942.
XX      PR      17-JUN-1998; 98US-0099041.
XX      PR      08-DEC-1998; 98US-0207359.
XX
XX      (MILL-) MILLENNIUM PHARM INC.
XX      PI
XX      Bertin J;
XX      WPI; 2002-391988/42.
XX
XX      Isolated Caspase Recruitment Domain (CARD) polypeptides (CARD-4L and
XX      PT      CARD-4S) useful for diagnosing and treating e.g. Parkinson's and
XX      PT      Alzheimer's disease, cancers and viral infections -
XX      PS
XX      Disclosure; Fig 18; 116pp; English.
XX
CC      The invention relates to novel isolated Caspase Recruitment Domain (CARD)
CC      polypeptides, CARD-4L and CARD-4S. The CARD proteins of the invention may
CC      be used to treat disorders associated with decreased CARD expression by
CC      supplementing the patient's own production of CARD. Disorders associated
CC      with the expression and activity of CARD include cancers (particularly
CC      follicular lymphomas, carcinomas associated with mutations in p53, and
CC      hormone-dependent tumours such as breast cancer, prostate cancer, and
CC      ovarian cancer), autoimmune disorders (such as systemic lupus
CC      erythematosus, immune-mediated glomerulonephritis), viral infections
CC      (such as those caused by herpes viruses, poxviruses, and adenoviruses),
CC      neurological diseases (such as Alzheimer's disease, Parkinson's
CC      disease, amyotrophic lateral sclerosis (ALS), retinitis pigmentosa,
CC      spinal muscular atrophy, and various forms of cerebellar degeneration),
CC      anaemia associated with chronic disease, aplastic anaemia, chronic
CC      neutropenia, and the myelodysplastic syndromes. This polynucleotide
CC      sequence represents the DNA of a human CARD relating to the invention.
XX      XX
XX      Sequence 32042 BP; 7389 A; 7540 C; 7721 G; 9392 T; 0 other;
SO

```

Query Match 22.6%; Score 27.4; DB 24; Length 32042;  
 Best Local Similarity 62.3%; Pred. No. 30;  
 Matches 43; Conservative 0; Mismatches 26; Indels 0; Gaps 0;

```

OY      28 TCGAGCGTCCCATGCTAATAATGACGCGCTCATGCTGCTGTTATTAAT 87
DB      18335 TTGACAGTACCCATCTCAATGAGTGTGAGAGTCTCTGCTATTTGCTTGG 18394
OY      88 GTTGATATA 96
DB      18395 GCTTTTAA 18403

```

## RESULT 18

ABO25024

ID ABO25024 standard; DNA: 628 BP.

XX ABO25024;

XX 12-JUL-2002 (first entry)

XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 11615.

XX Human: cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;  
 KW drug; side effect; cancer; central nervous system; cardiovascular;  
 KW gastrointestinal; respiratory system; single nucleotide polymorphism;  
 KW SNP; cell differentiation; ds.

XX Homo sapiens.

XX WO200218632-A2.

XX 07-MAR-2002.

XX 01-SEP-2001; 2001WO-EPI0074.

XX 01-SEP-2000; 2000DE-1043826.

XX 05-SEP-2000; 2000DE-1044543.

XX (EPIC-) EPIGENOMICS AG.

XX Olek A, Piepenbrock C, Berlin K, Guetig D;

XX WPI; 2002-371829/40.

XX Determining the degree of cytosine methylation in genomic DNA, useful  
 PT for diagnosis and prognosis, comprises selective hybridization of  
 PT amplicons from chemically treated DNA -

XX Claim 12; 56pp + Sequence Listing; 56pp; German.

XX This invention describes a novel method for determining the degree of  
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a  
 CC genomic sample of DNA. The sample is treated chemically to convert  
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic  
 CC DNA that contains the target C is amplified to form a labeled amplicon.  
 CC The amplicon is hybridised to two classes, each with at least one  
 CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers  
 CC and the degree of hybridisation to both classes is determined from the  
 CC label on the amplicon. From the ratio of labels hybridised to the two  
 CC classes of oligomers, the degree of methylation is calculated. The method  
 CC is used: (i) for diagnosis and/or prognosis of side effects of  
 CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders  
 CC of the central nervous, cardiovascular, gastrointestinal and respiratory  
 CC systems etc., particularly by detecting mutations or single nucleotide  
 CC polymorphisms (SNP's); and (ii) for differentiation of cell or tissue  
 CC types and for investigating cell differentiation. The method allows the  
 CC methylation status of many C residues to be determined simultaneously.  
 CC ABO13410-ABO54121 represent genomic DNA sequences used to illustrate the  
 CC method for determining the degree of cytosine methylation described in  
 CC the disclosure of the invention.

XX Sequence 628 BP; 161 A; 54 C; 160 G; 253 T; 0 other;

XX Query Match 22.3%; Score 27; DB 24; Length 628;

XX Best Local Similarity 60.0%; Pred. No. 11;

XX Matches 45; Conservative 0; Mismatches 30; Indels 0; Gaps 0;

OY 26 GATCGACGCTGCCATGCTGATTAATGACGGCTCCTGATCCATTTGCTGTGTTATTA 85

DB 509 GATCGACGGTGTATTTGTTAGTAATAGATGTTGATGATTCGTGCTGTTGTTATTTT 568

OY 86 ATGTTGTTAATTGA 100

DB 569 TAGTATTTGTTACGA 583

## RESULT 19

ABO25025/C

ID ABO25025 standard; DNA: 628 BP.

XX ABO25025;

XX 12-JUL-2002 (first entry)

XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 11616.

XX Human: cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;  
 KW drug; side effect; cancer; central nervous system; cardiovascular;  
 KW gastrointestinal; respiratory system; single nucleotide polymorphism;  
 KW SNP; cell differentiation; ds.

XX Homo sapiens.

XX WO200218632-A2.

XX 07-MAR-2002.

XX 01-SEP-2001; 2001WO-EPI0074.

XX 01-SEP-2000; 2000DE-1043826.

XX 05-SEP-2000; 2000DE-1044543.

XX (EPIC-) EPIGENOMICS AG.

XX Olek A, Piepenbrock C, Berlin K, Guetig D;

XX WPI; 2002-371829/40.

XX Determining the degree of cytosine methylation in genomic DNA, useful  
 PT for diagnosis and prognosis, comprises selective hybridization of  
 PT amplicons from chemically treated DNA -

XX Claim 12; 56pp + Sequence Listing; 56pp; German.

XX This invention describes a novel method for determining the degree of  
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a  
 CC genomic sample of DNA. The sample is treated chemically to convert  
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic  
 CC DNA that contains the target C is amplified to form a labeled amplicon.  
 CC The amplicon is hybridised to two classes, each with at least one  
 CC member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers  
 CC and the degree of hybridisation to both classes is determined from the  
 CC label on the amplicon. From the ratio of labels hybridised to the two  
 CC classes of oligomers, the degree of methylation is calculated. The method  
 CC is used: (i) for diagnosis and/or prognosis of side effects of  
 CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders  
 CC of the central nervous, cardiovascular, gastrointestinal and respiratory  
 CC systems etc., particularly by detecting mutations or single nucleotide  
 CC polymorphisms (SNP's); and (ii) for differentiation of cell or tissue  
 CC types and for investigating cell differentiation. The method allows the  
 CC methylation status of many C residues to be determined simultaneously.  
 CC ABO13410-ABO54121 represent genomic DNA sequences used to illustrate the  
 CC method for determining the degree of cytosine methylation described in  
 CC the disclosure of the invention.

XX Sequence 628 BP; 253 A; 160 C; 54 G; 161 T; 0 other;

XX Query Match 22.3%; Score 27; DB 24; Length 628;

XX Best Local Similarity 60.0%; Pred. No. 11;

XX Matches 45; Conservative 0; Mismatches 30; Indels 0; Gaps 0;

OY 26 GATCGACGCTGCCATGCTGATTAATGACGGCTCCTGATCCATTTGCTGTGTTATTA 85

DB 120 GATCGACGGTGTATTTGTTAGTAATAGATGTTGATGATTCGTGCTGTTGTTATTTT 61

OY 86 ATGTTGTTAATTGA 100

DB 60 TAGTATTTGTTACGA 46



AC	AAZ97380,
XX	
DT	18-APR-2000 (first entry)
XX	
DE	Human prostate cancer differentially expressed gene #241.
XX	
KW	Prostate cancer specific gene; cancer; tumour progression; diagnose;
KW	hyperproliferative cell growth; prostatic disorder; treatment;
KW	metastatic prostate cancer; benign prostate hyperplasia; BPH; ss.
XX	
OS	Homo sapiens.
XX	
PN	WO964594-A2.
XX	
PD	16-DEC-1999.
XX	
PF	10-JUN-1999; 99WO-US13181.
XX	
PR	11-JUN-1998; 98US-0088877.
PR	09-JUN-1999; 99US-0088877.
XX	
PA	(CHIR ) CHIRON CORP.
PI	Astel JH, Carroll E, Endege WO, Ford DM, Monahan JE, Schlegel R;
PI	Steinmann KE, Zhang J;
DR	WPI; 2000-116541/10.
XX	
PT	New isolated prostate cancer specific nucleic acids, used to develop
PT	products for the diagnosis and treatment of cancer -
XX	
PS	Claim 2; Page 173; 212pp; English.
XX	
CC	This sequence represents a prostate cancer specific nucleic acid
CC	sequence. The invention relates to a method for diagnosing cancer,
CC	tumour progression, hyperproliferative cell growth or accompanying
CC	biological and physical manifestations. The method involves contacting
CC	the biological sample with a probe that comprises a sequence capable of
CC	hybridising to any of the 339 nucleotide sequences given in the
CC	specification (see AAZ97140-297478) and detecting duplex formation. The
CC	products and methods of the invention can be used for the diagnosis,
CC	prognosis, and treatment of cancer, tumour progression,
CC	hyperproliferative cell growth, and accompanying physical and biological
CC	manifestations. They can be used particularly for prostatic disorders
CC	such as metastatic prostate cancer, localised prostate cancer, or benign
CC	prostate hyperplasia (BPH).
CC	
SQ	Sequence 486 BP; 137 A; 111 C; 128 G; 108 T; 2 other:
	Query Match            22.1%; Score 26.8; DB 21; Length 486;
	Best Local Similarity   59.0%; Pred. No. 12;
	Matches   46; Conservative   0; Mismatches   32; Indels   0; Gaps   0;
OY	33 GCTGCCCATGTGCATTAATGAGCGGTCCTGCATCATTGTGCTGTATATATGGTTGT 92
DB	303 GCCTTCTAAACTGCGGGAATTACAGTCTGTGAGCCACTGTGCTGTCAATAAATT 244
OY	93 ATAATTGAGCAGGACACA 110
DB	243 AATTCATCATCACATTACA 226
RESULT 23	
ID	AAZ97180/C
XX	AAZ97180 standard; CDNA; 1004 BP.
XX	
AC	AAZ97180;
XX	
DT	18-APR-2000 (first entry)
XX	
DE	Human prostate cancer differentially expressed gene #41.
XX	
KW	Prostate cancer specific gene; cancer; tumour progression; diagnose;

KM	hyperproliferative cell growth; prostatic disorder; treatment;
KM	metastatic prostate cancer; benign prostate hyperplasia; BPH; ss.
OS	Homo sapiens.
PN	MO964594-A2.
PD	16-DEC-1999.
PF	10-JUN-1999; 99WO-US3181.
PR	11-JUN-1998; 98US-0088877.
PR	09-JUN-1999; 99US-0088877.
PA	(CHIR ) CHIRON CORP.
PI	Astel JH, Carroll E, Endege WO, Ford DM, Monahan JE, Schlegel R;
PI	Steinmann KE, Zhang J;
DR	WPI; 2000-116541/10.
PT	New isolated prostate cancer specific nucleic acids, used to develop
PT	products for the diagnosis and treatment of cancer -
PS	Claim 2; Page 89; 212pp; English.
CC	This sequence represents a prostate cancer specific nucleic acid
CC	sequence. The invention relates to a method for diagnosing cancer,
CC	tumour progression, hyperproliferative cell growth or accompanying
CC	biological and physical manifestations. The method involves contacting
CC	the biological sample with a probe that comprises a sequence capable of
CC	hybridising to any of the 339 nucleotide sequences given in the
CC	specification (see AA297140-297478) and detecting duplex formation. The
CC	products and methods of the invention can be used for the diagnosis,
CC	prognosis and treatment of cancer, tumour progression,
CC	hyperproliferative cell growth, and accompanying physical and biological
CC	manifestations. They can be used particularly for prostatic disorders
CC	such as metastatic prostate cancer, localised prostate cancer, or benign
CC	prostate hyperplasia (BPH).
CC	xx
CC	xx
SO	Sequence 1004 BP: 266 A: 208 C: 218 G: 259 T: 53 other;
Query Match	22.1%; Score 26.8; DB 21; Length 1004;
Best Local Similarity	59.0%; Pred. No. 15;
Matches 46; Conservative 0; Mismatches 32; Indels 0; Gaps 0;	
OY	33 GCTGCCATGCGATAAATGACGGCTCCGATCCATGTGTCGTATTAATGTTGT 92
DB	299 GCTTCTTAACGCTGCGAATTACAGTCGTGACGCCACTGTGCTGTCTCAATAACTTTT 240
OY	93 ATAATTGACGACGACACA 110
DB	239 AAATTCATCACATTACA 222
RESULT 24	
AA03458	
ID	AA03458 standard; DNA; 2033 BP.
XX	
AC	AA03458;
XX	
DT	13-JUN-2001 (first entry)
XX	
DE	Zea mays modified ubiquitin regulatory system (mURS) DNA.
XX	
KM	Maize; modified ubiquitin regulatory system; mURS; heatshock element;
KW	transgenic plant; gene regulation; ds.
XX	
OS	Zea mays.
XX	
FT	Key Location/Qualifiers
FT	misc-feature 718..723
FT	/tag= a

```

FT      /label="KpnI-site"
FT      /note="Replaces the overlapping heatshock elements
XX      in the wild-type URS"
PN      WO200118220-A1.
XX
PD      15-MAR-2001.
XX
PF      07-SEP-2000; 2000WO-EP08690.
XX
PR      09-SEP-1999; 99EP-0307158.
XX
PA      (MONS ) MONSANTO PLC.
XX
PI      Goldsbrough A;
XX
DR      WPI; 2001-235203/24.
XX
PT      DNA sequence comprising a ubiquitin regulatory system, which lacks
PT      heatshock elements or is not heat inducible, useful for transforming
PT      plants (e.g. corn, wheat or barley) and regulating the expression of a
PT      structural gene under its control.
XX
PS      Claim 4; Fig 3; 39pp; English.
XX
CC      The present sequence is Zea mays modified ubiquitin regulatory system
CC      (mURS) DNA which lacks heatshock elements or is not heat inducible.
CC      The mURS DNA sequence is useful for transforming cells, particularly
CC      plant cells and creating transgenic plants. It is useful for regulating
CC      the expression of a structural gene, particularly when stressed with
CC      elevated temperature. The expression of associated DNA sequences, which
CC      is mediated by the present ubiquitin regulatory system, in transformed
CC      plant cells is stable and not influenced by environmental changes in
CC      temperature, which would normally affect expression mediated by a
CC      non-modified system.
XX
SQ      Sequence 2033 BP; 488 A; 424 C; 423 G; 698 T; 0 other:
XX
Query Match      22.1%; Score 26.8; DB 22; Length 2033;
Best Local Similarity 53.9%; Pred. No. 19;
Matches 55; Conservative 0; Mismatches 47; Indels 0; Gaps 0;
XX
OY      3 TTTCGCTAGCCTCGTGAGACATCGACGCTGCCATCGTAAATGACGCTCCTG 62
DB      1493 TATCGATCTAGCATGATGATACGTTGATCGCGGTTTACTGATGATATACAGAGATG 1552
OY      63 ATCCATTGTTGCTGTGTTATTAATGTTGTAATTAATGAGCAG 104
DB      1553 CTTTGTTCGCTGTGTTGATGATGCTGTGTTGGCGG 1594
XX
RESULT 25
AAZ99961
ID      AAZ99961 standard; DNA; 2037 BP.
XX
AC      AAZ99961;
XX
DT      25-JUL-2000 (first entry)
XX
DE      Nucleotide sequence of a cloned fragment of p97-U3.
XX
KM      Wheat; starch branching enzyme II; SBEII; SBEII-1; starch; SBEII-2;
KM      gelatinisation onset; transgenic plant; foodstuff; bakery product; ds.
XX
OS      Zea mays.
XX
PN      WO200015810-A1.
XX
PD      23-MAR-2000.
XX
PF      09-SEP-1999; 99WO-GB03011.
XX
PR      10-SEP-1998; 98EP-0307337.

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XX      (PLAN-) PLANT BREEDING INT CAMBRIDGE LTD.
PA      Goldsbrough A, COLLIVER S;
XX
DR      WPI; 2000-271446/23.
XX
PT      DNA encoding wheat starch branching enzyme II isoforms, useful in
PT      altering the characteristics of a plant, especially elevated starch
PT      gelatinization onset and/or peak temperature.
XX
PS      Disclosure; Fig 29; 197pp; English.
XX
CC      The present sequence represents a cloned fragment of p97-U3, which
CC      is used in the course of the invention. The specification describes novel
CC      subclasses of wheat starch branching enzyme II (SBEII), designated
CC      SBEII-1 and SBEII-2. The SBEII-1 genes are thought to have similar
CC      functional properties to the maize SBEIIb gene. Starch branching enzymes
CC      catalyse the formation of the alpha-1,6 linkages, creating branch points
CC      in the growing starch molecule, via hydrolysis of an alpha-1,4 linkage
CC      followed by reattachment of the released alpha-1,4-glucan chain to the
CC      same or another glucosyl chain. SBEII polypeptides can be used to alter
CC      the characteristics of a plant, in particular to alter starch so that
CC      it has an elevated gelatinisation onset and/or peak temperature. Starch
CC      obtained from transgenic plants is useful in the preparation or
CC      processing a foodstuff, particularly bakery products.
XX
SQ      Sequence 2037 BP; 488 A; 426 C; 423 G; 700 T; 0 other:
XX
Query Match      22.1%; Score 26.8; DB 21; Length 2037;
Best Local Similarity 53.9%; Pred. No. 19;
Matches 55; Conservative 0; Mismatches 47; Indels 0; Gaps 0;
XX
OY      3 TTTCGCTAGCCTCGTGAGACATCGACGCTGCCATCGTAAATGACGCTCCTG 62
DB      1495 TATCGATCTAGCATGATGATACATGTCATGCGGTTTACTGATGATATACAGAGATG 1554
OY      63 ATCCATTGTTGCTGTGTTATTAATGTTGTAATTAATGAGCAG 104
DB      1555 CTTTGTTCGCTGTGTTGATGATGCTGTGTTGGCGG 1596
XX
RESULT 26
AAH14585/C
ID      AAH14585 standard; cDNA; 2650 BP.
XX
AC      AAH14585;
XX
DT      26-JUN-2001 (first entry)
XX
DE      Human cDNA sequence SEQ ID NO:12185.
XX
KM      Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX
OS      Homo sapiens.
XX
PN      EP1074617-A2.
XX
PD      07-FEB-2001.
XX
PF      28-JUL-2000; 2000EP-0116126.
XX
PR      29-JUL-1999; 99JP-0248036.
PR      27-AUG-1999; 99JP-0300253.
PR      11-JAN-2000; 2000JP-0118776.
PR      02-MAY-2000; 2000JP-0183767.
PR      09-JUN-2000; 2000JP-0241899.
XX
PA      (HELI-) HELIX RES INST.
XX
PI      Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J,
PI      Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
XX

```



DR WPI: 2001-318749/34.  
 XX Primer sets for synthesizing polynucleotides, particularly the 5602  
 PT full-length cDNAs defined in the specification, and for the detection  
 PT and/or diagnosis of the abnormality of the proteins encoded by the  
 PT full-length cDNAs -  
 XX  
 PS Claim 8; SEQ ID 12185; 2537pp + CD ROM; English.  
 XX  
 CC The present invention describes primer sets for synthesizing 5602  
 CC full-length cDNAs defined in the specification. Where a primer set  
 CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary  
 CC to the complementary strand of a polynucleotide which comprises one of  
 CC the 5602 nucleotide sequences defined in the specification, where the  
 CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination  
 CC of an oligonucleotide comprising a sequence complementary to the  
 CC complementary strand of a polynucleotide which comprises a 5'-end  
 CC sequence and an oligonucleotide comprising a sequence complementary to a  
 CC polynucleotide which comprises a 3'-end sequence, where the  
 CC oligonucleotide comprises at least 15 nucleotides and the combination of  
 CC the 5'-end sequence/3'-end sequence is selected from those defined in  
 CC the specification. The primer sets can be used in antisense therapy and  
 CC in gene therapy. The primers are useful for synthesizing polynucleotides,  
 CC particularly full-length cDNAs. The primers are also useful for the  
 CC detection and/or diagnosis of the abnormality of the proteins encoded by  
 CC the full-length cDNAs. The primers allow obtaining of the full-length  
 CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and  
 CC AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to  
 CC AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632  
 CC represent oligonucleotides, all of which are used in the exemplification  
 CC of the present invention.  
 CC  
 XX  
 SQ Sequence 2650 BP; 849 A; 495 C; 607 G; 699 T; 0 other;  
 Query Match 22.1%; Score 26.8; DB 22; Length 2650;  
 Best Local Similarity 59.0%; Pred. No. 21;  
 Matches 46; Conservative 0; Mismatches 32; Indels 0; Gaps 0;  
 OY 33 GCTGCCATGCTGATTAATGAGCGTCTGATTCATTTGTTGTTATTAATGTTGT 92  
 DB 1858 GCTTCTTAACCTGCTGGAATTAACATGCTGAGCCACTGCTGCTCATTAACCTTT 1799  
 OY 93 ATAATTGAGCAGACACA 110  
 DB 1798 AATTCATCATTACATTACA 1781

RESULT 27  
 AAH16327/c  
 ID AAH16327 standard; cDNA; 2897 BP.  
 XX  
 AC AAH16327;  
 XX  
 DT 26-JUN-2001 (first entry)  
 XX  
 DE Human cDNA sequence SEQ ID NO:15232.  
 XX  
 DE Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN EP1074617-A2.  
 XX  
 PD 07-FEB-2001.  
 XX  
 PF 28-JUL-2000; 2000EP-0116126.  
 XX  
 PR 29-JUL-1999; 99JP-0248036.  
 PR 27-AUG-1999; 99JP-0300253.  
 PR 11-JAN-2000; 2000JP-0118776.  
 PR 02-MAY-2000; 2000JP-0183767.  
 PR 09-JUN-2000; 2000JP-0241899.  
 XX

PA (HELI-) HELIX RES INST.  
 XX  
 PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;  
 PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;  
 XX  
 DR WPI: 2001-318749/34.  
 XX  
 PS Claim 8; SEQ ID 15232; 2537pp + CD ROM; English.  
 XX  
 CC The present invention describes primer sets for synthesizing 5602  
 CC full-length cDNAs defined in the specification. Where a primer set  
 CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary  
 CC to the complementary strand of a polynucleotide which comprises one of  
 CC the 5602 nucleotide sequences defined in the specification, where the  
 CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination  
 CC of an oligonucleotide comprising a sequence complementary to the  
 CC complementary strand of a polynucleotide which comprises a 5'-end  
 CC sequence and an oligonucleotide comprising a sequence complementary to a  
 CC polynucleotide which comprises a 3'-end sequence, where the  
 CC oligonucleotide comprises at least 15 nucleotides and the combination of  
 CC the 5'-end sequence/3'-end sequence is selected from those defined in  
 CC the specification. The primer sets can be used in antisense therapy and  
 CC in gene therapy. The primers are useful for synthesizing polynucleotides,  
 CC particularly full-length cDNAs. The primers are also useful for the  
 CC detection and/or diagnosis of the abnormality of the proteins encoded by  
 CC the full-length cDNAs. The primers allow obtaining of the full-length  
 CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and  
 CC AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to  
 CC AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632  
 CC represent oligonucleotides, all of which are used in the exemplification  
 CC of the present invention.  
 CC  
 XX  
 SQ Sequence 2897 BP; 882 A; 564 C; 696 G; 755 T; 0 other;  
 Query Match 22.1%; Score 26.8; DB 22; Length 2897;  
 Best Local Similarity 59.0%; Pred. No. 22;  
 Matches 46; Conservative 0; Mismatches 32; Indels 0; Gaps 0;  
 OY 33 GCTGCCATGCTGATTAATGAGCGTCTGATTCATTTGTTGTTATTAATGTTGT 92  
 DB 2080 GCTTCTTAACCTGCTGGAATTAACATGCTGAGCCACTGCTGCTCATTAACCTTT 2021  
 OY 93 ATAATTGAGCAGACACA 110  
 DB 2020 AATTCATCATTACATTACA 2003

RESULT 28  
 AAT31290/c  
 ID AAT31290 standard; cDNA; 3095 BP.  
 XX  
 AC AAT31290;  
 XX  
 DT 24-FEB-1997 (first entry)  
 XX  
 DE Mouse poly-immunoglobulin receptor, cDNA.  
 XX  
 DE Mouse; immunoglobulin; receptor; protection protein; mutants;  
 KW heavy chain; antigen binding domain; protection; pathogen;  
 KW mucosal; environment; gastrointestinal; passive; immunisation;  
 KW Guy's 13 antibody; prevention; dental caries; Streptococcus;  
 KW poly; sorbinus; murine; ss.  
 XX  
 OS Mus musculus.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 85..2400  
 FT /tag= a

```

XX XX WO9621012-A1.
XX PN
XX PD 11-JUL-1996.
XX XX
PF 27-DEC-1995: 95WO-US16689.
XX XX
PR 04-MAY-1995: 95US-0434000.
PR 30-DEC-1994: 94US-0367395.
XX XX
PA (PLAN-) PLANT BIOTECHNOLOGY INC.
PA (UNNE-) UNITED MEDICAL & DENTAL SCHOOLS GUYS.
PA (PLAN-) PLANET BIOTECHNOLOGY INC.
PI Hiatt AC, Lehner T, Ma JKC;
XX XX
DR WPI: 1996-333987/33.
DR P-PsDB: AAM03180.
XX XX
PT Immunoglobulin and protection protein complex and its prodn. in
PT plants - useful for passive immunisation against mucosal antigens,
PT esp. against S. mutans and S. sorbinus to prevent dental caries
PS
XX Disclosure: Pages 117-121; 152pp; English.
XX
CC The present sequence encodes the mouse poly-immunoglobulin (Ig)
CC receptor, a portion of which corresp. to residues 1-627, pref:
CC 1-606, or esp. residues 13-45, 1-120, 110-230, 210-340, 320-450,
CC 440-550, 550-606 or 550-627 comprises a protection protein (PP).
CC The Ig of the invention comprises a PP as above in association with
CC an Ig derived heavy chain, having at least a portion of an antigen
CC (Ag) binding domain. The PP protects the Ig in harsh mucosal, e.g.
CC gastrointestinal, environments, therefore enhancing its
CC effectiveness in passively immunising animals against mucosal
CC pathogens. The Ag binding domain is specifically derived from the
CC Guy's 13 antibody, and the Ig can be used to prevent dental caries
CC by binding, e.g. Streptococcus mutans serotypes C, e and f, or
CC S. sorbinus serotypes d and g.
XX
SQ Sequence 3095 BP; 861 A; 796 C; 784 G; 654 T; 0 other;
Query Match 22.1%; Score 26.8; DB 17; Length 3095;
Best Local Similarity 64.5%; Pred. No. 22;
Matches 40; Conservative 0; Mismatches 22; Indels 0; Gaps 0;
OY 30 GACGCTCCCATGCTGATAAATGAGCGTCCGATTCATGTTGTTATTAATCT 89
ID 111 111 111 111 111 111 111 111 111 111 111 111 111
DB 760 GACCTTCTCCAGCTTGCGAGATGTACAGCCACCATCATTTGCTTGGTAGCTAATCT 701
OY 90 TG 91
ID 11
DB 700 TG 699

```

```

PF 23-MAR-2001; 2001WO-US09231.
XX XX
PR 23-MAR-2000; 2000US-191637P.
PR 11-JUL-2000; 2000US-0614150.
XX XX
PA (PEKE ) PE CORP NY.
XX XX
PI Venter JC, Adams M, Li PWD, Myers EW;
XX XX
DR WPI: 2001-656860/75.
XX XX
PT New isolated nucleic acid detection reagent for detecting 1000 or more
PT genes from Drosophila and for elucidating cell signalling and cell-cell
PT interactions -
XX XX
PS Claim 1; SEQ ID NO 1375; 21pp + Sequence Listing; English.
XX
CC The invention relates to an isolated nucleic acid detection reagent
CC capable of detecting 1000 or more genes from Drosophila. The invention is
CC useful in developmental biology and in elucidating cell signalling and
CC cell-cell interactions in higher eukaryotes for the development of
CC insecticides, therapeutics and pharmaceutical drugs. The invention
CC discloses genomic DNA sequences (AB16176-AB130511), expressed DNA
CC sequences (AB101840-AB16175) and the encoded proteins
CC (AB57737-AB72072).
CC The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pcl_sequences.
XX
SQ Sequence 13255 BP; 3879 A; 2652 C; 2622 G; 4102 T; 0 other;
Query Match 22.1%; Score 26.8; DB 23; Length 13255;
Best Local Similarity 61.4%; Pred. No. 37;
Matches 43; Conservative 0; Mismatches 27; Indels 0; Gaps 0;
OY 44 TGATAATGAGCGCTCGATTCATGTTGCTGTTATTAATGTTGTTATTAATGAGCA 103
ID 11111 111 111 111 111 111 111 111 111 111 111 111
DB 2315 TGATCTCGATTAATGATTAATTTTCGCTGTTCAATTCGTAAATGAGCA 2256
OY 104 GGACACACACA 113
ID 1111 111
DB 2255 GGACATATAAA 2246

```

```

RESULT 30
AAV15422/C
ID AAV15422 standard; DNA; 29392 BP.
XX
AC AAV15422;
XX
DT 11-JUN-1998 (first entry)
XX
DE Mouse poly Ig receptor protein gene.
XX
KM Mouse; poly Ig receptor protein; p1gR protein; p1g; deficiency;
KM knockout mouse; disease model; ds.
XX
OS Mus sp.
XX
PN JPI0057066-A.
XX
PD 03-MAR-1998.
XX
PF 19-AUG-1996; 96JP-0217154.
XX
PR 19-AUG-1996; 96JP-0217154.
XX
PA (HONS ) YAKULT HONSHA KK.
XX
DR WPI: 1998-254323/23.
XX
PT Mouse p1g receptor protein gene - used for preparing gene knockout
mice, useful for study of human poly Ig receptor protein deficiency

```

XX Claim 1: Page 4-14; 18pp; Japanese.  
XX  
XX The present sequence represents the mouse poly Ig receptor protein  
CC gene, which has a 29392 bp sequence. The new gene can be used to  
CC produce a gene knockout mouse, useful as a disease model of human  
CC poly Ig receptor protein deficiency.  
XX  
SQ Sequence 29392 BP; 8318 A; 6747 C; 6514 G; 7813 T; 0 other;  
  
Query Match 22.1%; Score 26.8; DB 19; Length 29392;  
Best Local Similarity 64.5%; Pred. No. 48;  
Matches 40; Conservative 0; Mismatches 22; Indels 0; Gaps 0;  
  
QY 30 GACGCTGCCATGCTGATTAATGAGCGGCTGATTCATTTGCTGTGTTATATGCT 89  
DB 21417 GACCTTCTCCACTGCTGCGATGATACGCCACATCATTTGCGTTAGTACTATATGT 21358  
QY 90 TG 91  
DB 21357 TG 21356  
  
RESULT 31  
AAK78746  
ID AAK78746 standard; DNA; 33513 BP.  
XX  
AC AAK78746;  
XX  
DT 07-NOV-2001 (first entry)  
XX  
DE Human immune/haematopoietic antigen genomic sequence SEQ ID NO:33558.  
XX  
KW Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;  
XX cytosolic; gene therapy; vaccine; metastasis; ds.  
OS Homo sapiens.  
XX  
XX WO200157182-A2.  
PN  
PD 09-AUG-2001.  
XX  
PF 17-JAN-2001; 2001WO-US01354.  
XX  
PR 31-JAN-2000; 2000US-0179065.  
PR 04-FEB-2000; 2000US-0180628.  
PR 24-FEB-2000; 2000US-0184664.  
PR 02-MAR-2000; 2000US-0186350.  
PR 16-MAR-2000; 2000US-0189874.  
PR 17-MAR-2000; 2000US-0190076.  
PR 18-APR-2000; 2000US-0198123.  
PR 19-MAY-2000; 2000US-0205515.  
PR 07-JUN-2000; 2000US-0209467.  
PR 28-JUN-2000; 2000US-0214886.  
PR 30-JUN-2000; 2000US-0215135.  
PR 07-JUL-2000; 2000US-0216647.  
PR 11-JUL-2000; 2000US-0216880.  
PR 11-JUL-2000; 2000US-0217487.  
PR 14-JUL-2000; 2000US-0217496.  
PR 14-JUL-2000; 2000US-0218290.  
PR 26-JUL-2000; 2000US-0220963.  
PR 26-JUL-2000; 2000US-0220964.  
PR 14-AUG-2000; 2000US-0224518.  
PR 14-AUG-2000; 2000US-0224519.  
PR 14-AUG-2000; 2000US-0225213.  
PR 14-AUG-2000; 2000US-0225214.  
PR 14-AUG-2000; 2000US-0225266.  
PR 14-AUG-2000; 2000US-0225267.  
PR 14-AUG-2000; 2000US-0225268.  
PR 14-AUG-2000; 2000US-0225270.  
PR 14-AUG-2000; 2000US-0225447.  
PR 14-AUG-2000; 2000US-0225757.  
PR 14-AUG-2000; 2000US-0225758.

PR 14-AUG-2000; 2000US-0225759.  
PR 18-AUG-2000; 2000US-0226279.  
PR 22-AUG-2000; 2000US-0226681.  
PR 22-AUG-2000; 2000US-0226868.  
PR 22-AUG-2000; 2000US-0227182.  
PR 23-AUG-2000; 2000US-0227009.  
PR 30-AUG-2000; 2000US-0228924.  
PR 01-SEP-2000; 2000US-0229287.  
PR 01-SEP-2000; 2000US-0229343.  
PR 01-SEP-2000; 2000US-0229344.  
PR 01-SEP-2000; 2000US-0229345.  
PR 05-SEP-2000; 2000US-0229509.  
PR 05-SEP-2000; 2000US-0229513.  
PR 06-SEP-2000; 2000US-0230437.  
PR 06-SEP-2000; 2000US-0230438.  
PR 08-SEP-2000; 2000US-0231242.  
PR 08-SEP-2000; 2000US-0231243.  
PR 08-SEP-2000; 2000US-0231244.  
PR 08-SEP-2000; 2000US-0231413.  
PR 08-SEP-2000; 2000US-0231414.  
PR 08-SEP-2000; 2000US-0232080.  
PR 08-SEP-2000; 2000US-0232081.  
PR 12-SEP-2000; 2000US-0231968.  
PR 14-SEP-2000; 2000US-0232397.  
PR 14-SEP-2000; 2000US-0232398.  
PR 14-SEP-2000; 2000US-0232399.  
PR 14-SEP-2000; 2000US-0232400.  
PR 14-SEP-2000; 2000US-0232401.  
PR 14-SEP-2000; 2000US-0233063.  
PR 14-SEP-2000; 2000US-0233064.  
PR 14-SEP-2000; 2000US-0233065.  
PR 21-SEP-2000; 2000US-0234223.  
PR 21-SEP-2000; 2000US-0234274.  
PR 25-SEP-2000; 2000US-0234997.  
PR 25-SEP-2000; 2000US-0234998.  
PR 26-SEP-2000; 2000US-0235484.  
PR 27-SEP-2000; 2000US-0235834.  
PR 27-SEP-2000; 2000US-0235836.  
PR 29-SEP-2000; 2000US-0236327.  
PR 29-SEP-2000; 2000US-0236367.  
PR 29-SEP-2000; 2000US-0236368.  
PR 29-SEP-2000; 2000US-0236369.  
PR 29-SEP-2000; 2000US-0236370.  
PR 02-OCT-2000; 2000US-0236802.  
PR 02-OCT-2000; 2000US-0237037.  
PR 02-OCT-2000; 2000US-0237038.  
PR 02-OCT-2000; 2000US-0237039.  
PR 02-OCT-2000; 2000US-0237040.  
PR 13-OCT-2000; 2000US-0239335.  
PR 13-OCT-2000; 2000US-0239337.  
PR 20-OCT-2000; 2000US-0240960.  
PR 20-OCT-2000; 2000US-0241221.  
PR 20-OCT-2000; 2000US-0241785.  
PR 20-OCT-2000; 2000US-0241786.  
PR 20-OCT-2000; 2000US-0241787.  
PR 20-OCT-2000; 2000US-0241808.  
PR 20-OCT-2000; 2000US-0241809.  
PR 20-OCT-2000; 2000US-0241826.  
PR 01-NOV-2000; 2000US-0244617.  
PR 08-NOV-2000; 2000US-0244674.  
PR 08-NOV-2000; 2000US-0246475.  
PR 08-NOV-2000; 2000US-0246476.  
PR 08-NOV-2000; 2000US-0246477.  
PR 08-NOV-2000; 2000US-0246478.  
PR 08-NOV-2000; 2000US-0246523.  
PR 08-NOV-2000; 2000US-0246524.  
PR 08-NOV-2000; 2000US-0246525.  
PR 08-NOV-2000; 2000US-0246526.  
PR 08-NOV-2000; 2000US-0246527.  
PR 08-NOV-2000; 2000US-0246528.  
PR 08-NOV-2000; 2000US-0246532.  
PR 08-NOV-2000; 2000US-0246609.  
PR 08-NOV-2000; 2000US-0246610.



```

XX  Human; beta-amyloid precursor protein; C-terminal peptide;
KW  gene transfer; transgenic animal; Alzheimer disease model;
XX  gene therapy; ss.
XX  Homo sapiens.
XX  EP653154-A.
XX  17-MAY-1995.
XX  07-NOV-1994; 94EP-0117512.
XX  12-NOV-1993; 93JP-0306026.
XX  (FARH ) HOECHST JAPAN LTD.
PA  (FARH ) HOECHST JAPAN KK.
XX  Kawarabayashi T, Kobayashi T, Sato M, Shoji M, Tada N;
XX  WPI: 1995-180492/24.
DR  P-PSDB; AAR74694.
XX  Transgenic animal model for Alzheimer's disease - contains DNA encoding
PT  part of beta-amyloid precursor protein in a gene construct designed for
XX  over-expression in various cell types
XX  Claim 2; Page 11; 32pp; English.
XX  The sequence encodes a human brain beta-amyloid precursor protein
CC  (APP) C-terminal peptide. The DNA may be transferred along with an
CC  APP signal peptide gene (e.g. AA088695) into somatic and germ cells of
CC  a non-human mammal, and the resulting transgenic animal may be used
CC  as a model for Alzheimer disease (AD). The animal model exhibits
CC  symptoms similar to AD, producing large quantities of APP C-terminal
CC  peptide, death of neuron cells in pyramidal cells at cerebral
CC  amyloid regions, increases in glial cells and deposition of
CC  abnormally phosphorylated tau protein. The animal model may be
CC  used to develop new therapies for AD, including gene therapy
CC  strategies.
XX  Sequence 297 BP; 86 A; 64 C; 78 G; 69 T; 0 other;
SO
XX
Query Match 22.0%; Score 26.6; DB 16; Length 297;
Best Local Similarity 58.0%; Pred. No. 12;
Matches 47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;
QY 11 TAGCCTGGTGGACAGATCGACGCTGCCCATCGATTAATGAGCGTCCATGCATTG 70
DB 122 TAGCGACAGTATGCTCATCTGATGCTGAAGAGAACAGTACACATCCATTG 181
QY 71 TTGGTTGTATTAAATGTTG 91
DB 182 ATCATGGTGTGTGGAGGTTG 202

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XX  17-MAY-1995.
XX  07-NOV-1994; 94EP-0117512.
XX  12-NOV-1993; 93JP-0306026.
XX  (FARH ) HOECHST JAPAN LTD.
PA  (FARH ) HOECHST JAPAN KK.
XX  Kawarabayashi T, Kobayashi T, Sato M, Shoji M, Tada N;
XX  WPI: 1995-180492/24.
DR  P-PSDB; AAR74695.
XX  Transgenic animal model for Alzheimer's disease - contains DNA encoding
PT  part of beta-amyloid precursor protein in a gene construct designed for
XX  over-expression in various cell types
XX  Claim 2; Page 12-13; 32pp; English.
XX  The sequence encodes a human brain beta-amyloid precursor protein
CC  (APP) mutant C-terminal peptide, and differs from AA088696 by a Glu
CC  to Gln conversion at codon 22. The DNA may be transferred along
CC  with an APP signal peptide gene (e.g. AA088695) into somatic and germ
CC  cells of a non-human mammal, and the resulting transgenic animal may
CC  be used as a model for Alzheimer disease (AD). The animal model
CC  exhibits symptoms similar to AD, producing large quantities of APP
CC  C-terminal peptide, death of neuron cells in pyramidal cells at
CC  cerebral amyloid regions, increases in glial cells and deposition
CC  of abnormally phosphorylated tau protein. The animal model may
CC  be used to develop new therapies for AD, including gene therapy
CC  strategies.
XX  Sequence 297 BP; 86 A; 65 C; 77 G; 69 T; 0 other;
SO
XX
Query Match 22.0%; Score 26.6; DB 16; Length 297;
Best Local Similarity 58.0%; Pred. No. 12;
Matches 47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;
QY 11 TAGCCTGGTGGACAGATCGACGCTGCCCATCGATTAATGAGCGTCCATGCATTG 70
DB 122 TAGCGACAGTATGCTCATCTGATGCTGAAGAGAACAGTACACATCCATTG 181
QY 71 TTGGTTGTATTAAATGTTG 91
DB 182 ATCATGGTGTGTGGAGGTTG 202

```

```

RESULT 35
AA042665
ID AA042665 standard; cDNA; 300 BP.
XX
AC AA042665;
XX
DE 27-OCT-1993 (first entry)
XX
DE Full-length beta-amyloid protein coding region.
XX
KW Alzheimer's Disease; Amyloid Plaque Core Protein; APCP;
KW neuritic plaque; ds.
XX
XX Homo sapiens.
XX
FH Key location/qualifiers
FT CDS 1..300
FT /*tag= a
FT /product= beta-amyloid_protein
FT /note= "full-length coding sequence obtained by
FT combining overlapping clones SM2W3 and
FT SM2W4 (AA042664 and AA042663, respectively)"
FT misc_difference 160..162
FT /*tag= b

```

	/codon= seq; AAC; aa: Lys
	/note= "this codon is AAG in AAQ42664"
FT	
XX	
PN	US5320013-A.
PD	15-JUN-1993.
PF	17-NOV-1986;
PR	86US-0932193.
PR	17-NOV-1986;
PR	86US-0932193.
PR	31-DEC-1986;
PR	86US-0948376.
PR	30-JAN-1987;
PR	87US-0008810.
PR	18-AUG-1987;
PR	87US-0087002.
PR	30-NOV-1989;
PA	89US-0044118.
XX	(SCIO-) SCIOS NOVA INC.
PI	Cordell B, Ponte PA:
DR	WPI: 1993-205383/25.
DR	P-PSDB; AAR37866.
PT	DNA sequence useful for detection of Alzheimer's disease - for encoding beta amyloid core protein
XX	
PS	Disclosure: Fig 5: 40pp; English.
CC	A clone was obtained from the genomic library described in Lawn et al., Cell, 15:1157-1174 (1978) which included a 57 base pair segment which encodes amino acids 1-18 of the beta-amyloid protein, immediately preceded by a Methionine. A HindIII/XbaI fragment derived from the genomic clone and containing the 57bp segment was used to isolate cDNA fragments from a library prepared from temporal and parietal cortical tissue from a normal human brain. Lambda clone SM2w3 (AAQ42664) contains a 5' region segment which has a 6bp overlap with the 3' end of clone SM2w4 (AAQ42663). The full-length beta-amyloid protein coding sequence (AAQ42665), including an initiator Met residue which is probably processed in vivo, was obtained by combining the sequences of the two overlapping clones.
SO	Sequence 300 BP; 87 A; 65 C; 78 G; 70 T; 0 other;
Query Match	22.0%; Score 26.6; DB 14; Length 300;
Best Local Similarity	58.0%; Pred. No. 12;
Matches	47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;
OY	11 TAGCTCGGTGACAGATCAGCGTGCCTCATTAATAATGAGCGTCCTGATCCATTG 70 
Db	125 TAGCAGCAAGTAGTGCTCATCACCTTTGGTGATGCTGAACAAGAAGACATCATTC 184
OY	71 TTTCGTTGCTTATTATATGTTG 91 
Db	185 ATCATGTGTGTGTGGAGGTTG 205
RESULT 36	
AAK93386 ID	AAK93386 standard; DNA: 300 BP.
XX	
AC	AAK93386;
DT	27-JUN-2002 (first entry)
DE	DNA of wild type C99 portion of human APP.
XX	
KM	Neuroprotective; nootropic; transgenic fly; Alzheimer's disease; Abeta;
KM	Amyloid precursor protein; tissue-specific expression control; human APP;
KM	APP pathway modulator; gene therapy; ds.
OS	Homo sapiens.
XX	
WM	WO200226820-A2.
XX	

PD	04-APR-2002.
XX	
PF	01-OCT-2001; 2001WO-EP11345.
XX	
PR	29-SEP-2000; 2000US-236893P.
XX	
PR	14-JUN-2001; 2001US-298309P.
XX	
PA	(NOVS ) NOVARTIS AG.
XX	
PI	Cohen D, Dengler UJ, Finelli AL, Freuler F, Konsolaki M,
PI	Reinhardt MMH, Zusman S;
DR	WPI: 2002-315796/35.
XX	
PT	New transgenic fly, containing DNA encoding an Abeta portion of human
PT	APP, useful for identifying agents which modulate the APP pathway and
PT	which can be used to treat Alzheimer's disease .
XX	
P5	Claim 4; Page 84; 129pp; English.
XX	
CC	The invention relates to a transgenic fly whose genome comprises DNA
CC	encoding a polypeptide having the Abeta portion of human amyloid
CC	precursor protein (APP), fused to a signal sequence. The DNA sequence
CC	encodes a 123 (Abeta40) or 129 (Abeta42) amino acid sequence, given in
CC	the specification. The DNA sequence is operably linked to a tissue-
CC	specific expression control sequence. Expression of the sequence gives
CC	the fly an altered phenotype. The purpose of the invention is for
CC	identifying agents that inhibit or promote the expression and/or function
CC	of genes or encoded polypeptides which modify the APP pathway. The agent
CC	is a compound, triple helix DNA, antisense oligonucleotide, double
CC	stranded RNA molecule, ribozyme, or particularly an antibody. It is used
CC	to treat conditions such as Alzheimer's disease. The agent can be used as
CC	an APP pathway modulator or in gene therapy. This polynucleotide sequence
CC	represents the DNA of the wild type C99 portion of human APP of the
CC	invention.
SO	Sequence 300 BP; 87 A; 65 C; 79 G; 69 T; 0 other:
Query Match	22.0%; Score 26.6; DB 24; Length 300;
Best Local Similarity	58.0%; Pred. No. 12;
Matches 47; Conservative	0; Mismatches 34; Indels 0; Gaps 0
OY	11 TAGCGTGGGAGACAGATGCAGCCTCCCATGTCTATTAATGAGCGTCCATCAATTG 70
DB	122 TAGGGAAGGATCGTCATCACCTTGCTGATCTGAAGAACAAGTACATCAATTC 181
OY	71 TTCCTGTGTTAATTAAGTTG 91
DB	182 ATCATGCTGTGTGAGAGTTG 202
RESULT 37	
AAA339488	
ID	AAA39488 standard; DNA; 303 BP.
XX	
AC	AAA39488;
DT	24-AUG-2000 (first entry)
XX	
DE	Human APP C100 DNA fragment.
XX	
KW	APP; amyloid precursor protein; C100; transgene; Alzheimer's disease;
KW	gamma-secretase; alpha-secretase; ss.
OS	Homo sapiens.
XX	
PN	DEL9849073-A1.
FD	27-APR-2000.
XX	
PF	24-OCT-1998; 98DE-1049073.
XX	

PR 24-OCT-1998; 98DE-1049073.  
 XX (AVET ) AVENTIS PHARMA DEUT GMBH.  
 XX  
 PI Peraus G, Hoppe E, Baumeister R;  
 XX  
 DR WPI; 2000-304753/27.  
 XX  
 PT Transgene encoding amyloid precursor protein or its fragment, used to  
 PT produce transgenic nematodes used, e.g. to screen for agents for  
 PT treating Alzheimer's disease  
 PR  
 XX  
 PS Claim 3; Page 6; 16pp; German.  
 XX  
 CC This invention describes a novel transgene (I) which comprises (i) a  
 CC sequence (Ia) that encodes either the complete amyloid precursor protein  
 CC (APP) or a part of its Abeta peptide (Abp), but is not identical with  
 CC the part of the APP sequence that encodes the complete Abp; (ii)  
 CC optionally one or more additional coding and/or non-coding sequences and  
 CC (iii) a promoter functional in *Caenorhabditis elegans*. Cleavage of APP  
 CC by alpha-secretase does not lead to Abp, but cleavage with beta or  
 CC gamma-secretases does, so modulation of these activities is used to  
 CC reduce the amount of Abp, a major component of the plaque found in  
 CC Alzheimer patients, produced. (I), or expression vectors containing it,  
 CC are used to produce transgenic cells (optionally part of a non-human  
 CC animal), specifically *C. elegans*. The transgenic animals are used (i) to  
 CC identify gamma and/or alpha-secretase activities in *C. elegans*; (ii) to  
 CC identify/or characterize substances that inhibit gamma-secretase or  
 CC increase alpha-secretase activities, or (iii) to identify and/or  
 CC characterize agents potentially useful for treatment or prevention of  
 CC Alzheimer's disease. This sequence represents the human APP protein C100  
 CC fragment which is used in the construction of the transgene described in  
 CC the method of the invention.  
 CC  
 XX  
 SQ Sequence 303 BP; 87 A; 65 C; 80 G; 71 T; 0 other;  
 OY  
 Query Match 22.0%; Score 26.6; DB 21; Length 303;  
 Best Local Similarity 58.0%; Pred. No. 12;  
 Matches 47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;  
 OY 11 TAGCCTCGGTGACAGATCGACGCTGCCATCGTATTAATGAGCGTCCATGTCATTG 70  
 DB 125 TAGCGACAGTATGCTCATCTGATCGTGTGATGCTGAAGAAGAACAGTACACATCATTC 184  
 OY 71 TTGGTTGTTATTATATGTTG 91  
 DB 185 ATCATGTGTGTGTGAGAGTTG 205  
 DB  
 RESULT 38  
 AA088699  
 ID AA088699 standard; cDNA to mRNA; 309 BP.  
 XX  
 AC AA088699;  
 XX  
 DT 11-NOV-1995 (first entry)  
 XX  
 DE Beta-amyloid precursor protein C-terminal peptide gene.  
 XX  
 KW Human; beta-amyloid precursor protein; C-terminal peptide;  
 KW gene transfer; transgenic animal; Alzheimer disease model;  
 KW gene therapy; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN EP653154-A.  
 XX  
 PD 17-MAY-1995.  
 XX  
 PF 07-NOV-1994; 94EP-0117512.  
 XX  
 PR 12-NOV-1993; 93JP-0306026.  
 XX

PA (FARH ) HOECHST JAPAN LTD.  
 PA (FARH ) HOECHST JAPAN KK.  
 XX  
 PI Kawarabayashi T, Kobayashi T, Sato M, Shoji M, Tada N;  
 XX  
 DR WPI; 1995-180492/24.  
 DR P-PSDB; AAR74697.  
 XX  
 PT Transgenic animal model for Alzheimer's disease - contains DNA encoding  
 PT part of beta-amyloid precursor protein in a gene construct designed for  
 PT over-expression in various cell types  
 PR  
 XX  
 PS Claim 2; Page 15; 32pp; English.  
 XX  
 CC The sequence encodes a human brain beta-amyloid precursor protein  
 CC (APP) C-terminal peptide. The DNA may be transferred along with an  
 CC APP signal peptide gene (e.g. AA088695) into somatic and germ cells  
 CC of a non-human mammal, and the resulting transgenic animal may be  
 CC used as a model for Alzheimer disease (AD). The animal model  
 CC exhibits symptoms similar to AD, producing large quantities of APP  
 CC C-terminal peptide, death of neuron cells in pyramidal cells at  
 CC cerebral amyloid regions, increases in glial cells and deposition  
 CC of abnormally phosphorylated tau protein. The animal model may  
 CC be used to develop new therapies for AD, including gene therapy  
 CC strategies.  
 CC  
 XX  
 SQ Sequence 309 BP; 91 A; 64 C; 83 G; 71 T; 0 other;  
 OY  
 Query Match 22.0%; Score 26.6; DB 16; Length 309;  
 Best Local Similarity 58.0%; Pred. No. 12;  
 Matches 47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;  
 OY 11 TAGCCTCGGTGACAGATCGACGCTGCCATCGTATTAATGAGCGTCCATGTCATTG 70  
 DB 134 TAGCGACAGTATGCTCATCTGATCGTGTGATGCTGAAGAAGAACAGTACACATCATTC 193  
 OY 71 TTGGTTGTTATTATATGTTG 91  
 DB 194 ATCATGTGTGTGTGAGAGTTG 214  
 DB  
 RESULT 39  
 AA088700  
 ID AA088700 standard; cDNA to mRNA; 309 BP.  
 XX  
 AC AA088700;  
 XX  
 DT 11-NOV-1995 (first entry)  
 XX  
 DE Beta-amyloid precursor protein C-terminal peptide mutant gene.  
 XX  
 KW Human; beta-amyloid precursor protein mutant; C-terminal peptide;  
 KW gene transfer; transgenic animal; Alzheimer disease model;  
 KW gene therapy; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN EP653154-A.  
 XX  
 PD 17-MAY-1995.  
 XX  
 PF 07-NOV-1994; 94EP-0117512.  
 XX  
 PR 12-NOV-1993; 93JP-0306026.  
 XX  
 PA (FARH ) HOECHST JAPAN LTD.  
 PA (FARH ) HOECHST JAPAN KK.  
 XX  
 PI Kawarabayashi T, Kobayashi T, Sato M, Shoji M, Tada N;  
 XX  
 DR WPI; 1995-180492/24.  
 DR P-PSDB; AAR74698.  
 XX

PT Transgenic animal model for Alzheimer's disease - contains DNA encoding  
 PT part of beta-amyloid precursor protein in a gene construct designed for  
 PT over-expression in various cell types  
 XX  
 XX  
 PS Claim 2: Page 16-17; 32pp; English.  
 CC The sequence encodes a human brain beta-amyloid precursor protein  
 CC (APP) mutant C-terminal peptide, and differs from AAO88699 by  
 CC conversion of Lys to Asn at codon-3 and Met to Leu at codon-4. The  
 CC DNA may be transferred along with an APP signal peptide gene (e.g.  
 CC AAO88699) into somatic and germ cells of a non-human mammal, and the  
 CC resulting transgenic animal may be used as a model for Alzheimer  
 CC disease (AD). The animal model exhibits symptoms similar to AD,  
 CC producing large quantities of APP C-terminal peptide, death of  
 CC neuron cells in pyramidal cells at cerebral amyloid regions,  
 CC increases in glial cells and deposition of abnormally  
 CC phosphorylated tau protein. The animal model may be used to develop  
 CC new therapies for AD, including gene therapy strategies.  
 CC  
 SQ Sequence 309 BP; 90 A; 65 C; 82 G; 72 T; 0 other;  
 Query Match 22.0%; Score 26.6; DB 16; Length 309;  
 Best Local Similarity 58.0%; Pred. No. 12;  
 Matches 47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;  
 QY 11 TAGCCTCGGTGACAGATCGAGCGTCCCGCTGATTAATGAGCGGCTCATTCATTG 70  
 DB 134 TAGCGAGAGTGCATCGTCACTGCTGATGCTGAAGAAGAACATACATCATTC 193  
 OY 71 TTGCTGTGTTATTAATGTTG 91  
 DB 194 ATCATGCTGTGCTGAGCGTTG 214  
 RESULT 40  
 AAD20982  
 ID AAD20982 standard; DNA; 315 BP.  
 AC AAD20982;  
 XX  
 DT 15-JAN-2002 (first entry)  
 XX  
 DE Human recombinant beta-amyloid precursor protein (betaAPP) C-83 DNA.  
 XX  
 KM Human; Alzheimer's disease; gamma-secretase; integral-membrane protein;  
 KM beta-amyloid precursor protein; betaAPP; ds.  
 XX  
 OS Homo sapiens.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 1..315  
 FT /\*tag= a  
 FT /product= "Human recombinant betaAPP protein (C-83)"  
 PN WO200175435-A2.  
 XX  
 PD 11-OCT-2001.  
 PD  
 PE 30-MAR-2001; 2001WO-US10453.  
 PE  
 PR 03-APR-2000; 2000US-194495P.  
 PR  
 XX (BRIM ) BRISTOL-MYERS SQUIBB CO.  
 PA  
 PI Roberts SB, Hendrick JP, Vintersky A, Lewis M, Smith DW, Pak R;  
 XX  
 DR WPI: 2001-648575/74.  
 DR P-PSDB: AAE12897.  
 XX  
 PT Novel gamma secretase protein, useful in the production of amyloids, is  
 PT capable of cleaving beta-amyloid precursor protein to produce beta  
 PT amyloid peptide  
 XX

PS Example 1: Fig 3; 127pp; English.  
 XX  
 CC The invention relates to the field of plaque amyloid deposits that are  
 CC the hallmarks of Alzheimer's disease. In particular, the invention  
 CC relates to an isolated, functionally-active protein that has  
 CC gamma-secretase activity. Gamma-secretase activity is necessary for  
 CC amyloid production. The present invention also relates to methods for  
 CC isolating integral-membrane proteins and protein complexes, including  
 CC the gamma-secretase protein of the invention. The method is useful for  
 CC monitoring the cleavage of beta-amyloid precursor protein (betaAPP)  
 CC by gamma-secretase. The present sequence is a DNA encoding human  
 CC recombinant betaAPP protein (C-83).  
 CC  
 SQ Sequence 315 BP; 75 A; 75 C; 92 G; 73 T; 0 other;  
 Query Match 22.0%; Score 26.6; DB 22; Length 315;  
 Best Local Similarity 58.0%; Pred. No. 12;  
 Matches 47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;  
 QY 11 TAGCCTCGGTGACAGATCGAGCGTCCCGCTGATTAATGAGCGGCTCATTCATTG 70  
 DB 137 TAGCGAGAGTGCATCGTCACTGCTGATGCTGAAGAAGAACATACATCATTC 196  
 OY 71 TTGCTGTGTTATTAATGTTG 91  
 DB 197 ATCATGCTGTGCTGAGCGTTG 217  
 RESULT 41  
 AAD23935  
 ID AAD23935 standard; DNA; 327 BP.  
 AC AAD23935;  
 XX  
 DT 12-MAR-2002 (first entry)  
 XX  
 DE Gamma-secretase substrate #1 DNA.  
 XX  
 KM Gamma-secretase substrate; beta-CTF domain; amyloid precursor protein;  
 KM APP; beta-secretase; Alzheimer's disease; fusion protein; ds.  
 XX  
 OS Unidentified.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 1..327  
 FT /\*tag= a  
 FT /product= "Gamma-secretase substrate"  
 FT /note= "This substrate contains APP beta-CTF domain and  
 FT hydrophilic moiety fusion"  
 FT misc-feature 1..300  
 FT /\*tag= b  
 FT /note= "APP beta-CTF domain DNA"  
 FT misc-feature 301..324  
 FT /\*tag= c  
 FT /note= "Region encoding hydrophilic moiety"  
 PN WO200183811-A1.  
 XX  
 PD 08-NOV-2001.  
 PD  
 PE 25-APR-2001; 2001WO-US13332.  
 PE  
 PR 01-MAY-2000; 2000US-201053P.  
 PR  
 XX (MERI ) MERCK & CO INC.  
 PA  
 PI Li Y, Xu M, Huang Q, Gardell S;  
 XX  
 DR WPI: 2002-066536/09.  
 DR P-PSDB: AAE14379.  
 XX  
 PT Novel gamma secretase substrate for assaying gamma secretase activity  
 PT and identifying compounds that regulate gamma secretase activity, e.g.  
 PT



PT Inhibitors of gamma secretase useful for treating Alzheimer's disease  
 XX  
 XX  
 XX  
 PS  
 XX Example 1: Page 12-13; 36pp; English.  
 CC The invention relates to gamma-secretase substrates containing  
 CC a hydrophilic polypeptide moiety covalently joined to the carboxyl  
 CC terminus of a beta-CTF domain. A beta-CTF domain is a polypeptide that  
 CC can be cleaved by gamma-secretase, and that approximates the  
 CC C-terminal fragment (amino acids 596-695) of amyloid precursor protein  
 CC (APP) produced after cleavage of APP by beta-secretase. The  
 CC hydrophilic polypeptide moiety increases the solubility of the  
 CC substrate in a zwitterionic detergent. The gamma-secretase substrate  
 CC is used in vitro assays employing zwitterionic detergent for  
 CC measuring gamma-secretase activity. The assay methods are useful for  
 CC purifying and characterizing the enzyme, to screen for compounds that  
 CC modulate gamma-secretase activity, and to test the ability of a  
 CC particular compound that affect gamma-secretase activity. The compounds  
 CC that modulate gamma-secretase activity include gamma-secretase  
 CC inhibitors which are useful in the treatment of Alzheimer's disease,  
 CC and in the characterisation of the biological importance of  
 CC gamma-secretase. The present sequence is a DNA encoding gamma-secretase  
 CC substrate of the invention. The substrate is a fusion protein  
 CC containing APP beta-CTF domain and a hydrophilic moiety.  
 XX  
 SO Sequence 327 BP; 100 A; 68 C; 85 G; 74 T; 0 other;  
 Query Match 22.0%; Score 26.6; DB 24; Length 327;  
 Best Local Similarity 58.0%; Pred. No. 12;  
 Matches 47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;  
 OY 11 TAGCCTCGGTGACAGATGACGCTGCCATGCTGATTAATGACGCTTCATTCATTG 70  
 DB 125 TAGCAGACAGTATCTCATACCTTGATGATGCTGAGAGAAGAACAGTACATCATTC 184  
 OY 71 TTCGTTGTTATTAATGTTG 91  
 DB 185 ATCATGTTGTGCTGGAGGTTG 205  
 DB 10-OCT-1996 (first entry)  
 XX  
 DE Familial Alzheimer's disease APP isoform 751 gene fragment.  
 XX  
 KW APP; amyloid precursor protein; isoform 751; inherent; familial;  
 KW Alzheimer's disease; mutation; diagnosis; transgenic model; study;  
 XX cognitive; beta A4 domain; exon 17; senility; ss.  
 OS Homo sapiens.  
 XX  
 XX Key Location/Qualifiers  
 FT Exon 1..336  
 FT /tag= a  
 FT /number= 17  
 FT /transl\_except= pos: 175-177, aa: Ile  
 FT /note= "encodes amino acids 640-751 of full APP  
 FT isoform 751, the translation exception  
 FT at posn. 175-177 is the site of a Val to  
 FT Ile mutation in isoform 751, bases 175-177  
 FT probably should be ATC and not TTC"  
 FT misc\_feature 43..168  
 FT /tag= b  
 FT /note= "encodes the beta-A4 domain"  
 PN W09606927-A1.  
 XX MO9606927-A1.  
 PD 07-MAR-1996.

XX  
 PF 28-AUG-1995; 95WO-US10920.  
 XX  
 PR 01-SEP-1994; 94US-0299872.  
 XX  
 XX (MERI ) MERCK & CO INC.  
 PA  
 PI Chen HY, Heavens RP, Singh G, Srinathsinghji DJS;  
 PI Smith DW, Trumbauer ME, Van Der Ploeg LHT, Vongs A,  
 PI Zheng H;  
 XX  
 DR WPI; 1996-160358/16.  
 DR P-PSDB; AAR93556.  
 XX  
 PT Transgenic animal expressing familial form of human amyloid  
 PT precursor protein used to evaluate compounds affecting  
 PT Alzheimer's disease and other cognitive disorders  
 PS  
 XX Example 1: Fig 7; 32pp; English.  
 CC  
 CC AAT18082 represents exon 17 of the amyloid precursor protein (APP)  
 CC isoform 751 gene from a patient diagnosed with familial Alzheimer's  
 CC disease (FAD). The sequence given corresponds to the coding sequence  
 CC for amino acids 640-751 of FAD APP 751. A feature of FAD is a Val to  
 CC Ile substitution at posn. 698 of the full APP (encoded by bases 175  
 CC to 177 of this sequence). DNA encoding this sequence was used to  
 CC construct expression vectors for the prodn. of transgenic animals (esp.  
 CC mice) carrying the FAD APP 751 mutation. The transgenic animals are  
 CC useful for the evaluation of test cpts. affecting Alzheimer's disease  
 CC and other cognitive disorders and for identification of new targets  
 CC in Alzheimer's disease since the progression of the disease can be  
 CC followed gradually.  
 CC N.B. the V-I mutation encoded at base 175-177 is given in the  
 CC specification as a TTC codon (most probably this should be ATC).  
 XX  
 SO Sequence 336 BP; 102 A; 69 C; 88 G; 77 T; 0 other;  
 Query Match 22.0%; Score 26.6; DB 17; Length 336;  
 Best Local Similarity 58.0%; Pred. No. 12;  
 Matches 47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;  
 OY 11 TAGCCTCGGTGACAGATGACGCTGCCATGCTGATTAATGACGCTTCATTCATTG 70  
 DB 161 TAGCAGACAGTATCTCATACCTTGATGATGCTGAGAGAAGAACAGTACATCATTC 220  
 OY 71 TTCGTTGTTATTAATGTTG 91  
 DB 221 ATCATGTTGTGCTGGAGGTTG 241  
 DB 16-MAR-1994 (first entry)  
 XX  
 DE Human brain Expressed Sequence Tag EST00434.  
 XX  
 KW Gene transcription product; genetic markers; tagging; in vivo;  
 KW Transcription; mapping; locations; chromosomes; chromosomal; ss.  
 OS Homo sapiens.  
 PN W09316178-A.  
 PF W09316178-A.  
 PD 19-AUG-1993.  
 XX  
 PF 12-FEB-1993; 93WO-US01294.  
 XX  
 PR 12-FEB-1992; 92US-0837195.  
 XX

PA (USSH ) US DEPT HEALTH & HUMAN SERVICE.  
 XX  
 PI Adams MD, Moreno RF, Venter CJ.  
 XX  
 DR WPI: 1993-272882/34.  
 XX  
 PT Enriched oligonucleotides and corresp. sequences - used as  
 PT markers for human genes transcribed in-vivo, facilitate tagging  
 PT of most human genes  
 XX  
 PS Example 4; Page 171; 500pp; English.  
 XX  
 CC The Expressed Sequence Tag was isolated from a human brain cDNA  
 CC library as part of a large set of ESTs which can be used as markers  
 CC for human genes transcribed in vivo. They can be used to facilitate  
 CC tagging of most human genes, for mapping locations of expressed genes  
 CC on chromosomes, for individual or forensic identification, for mapping  
 CC locations of disease-associated genes, for identification of tissue  
 CC type, and for prepn. of antisense sequences, probes and constructs.  
 CC EST00434 has a "poor" coding probability as evaluated using the  
 CC coding-region prediction program CRM. See also AA059041-061440.  
 CC  
 SQ Sequence 337 BP; 95 A; 63 C; 66 G; 107 T; 6 other;  
 XX  
 Query Match 22.0%; Score 26.6; DB 14; Length 337;  
 Best Local Similarity 54.9%; Pred. No. 12;  
 Matches 50; Conservative 0; Mismatches 41; Indels 0; Gaps 0;  
 OY 11 TAGCCTCGGTGACAGATCGATGCTGATTAATGAGCGTCCCTGATCCATTG 70  
 DB 58 TTAGCTTAGGCTTTCTGAGATCACTTCTCTCTTATTAATGAGAGATTAATCCATTG 117  
 OY 71 TTGCTGTGTATTATTAATGTTATTAATGAG 101  
 DB 118 CTCATTGAGTTGTTAATNAGACATAAATGAG 148  
 XX  
 RESULT 44  
 AAV07188  
 ID AAV07188 standard; DNA; 354 BP.  
 XX  
 AC AAV07188;  
 XX  
 DT 09-SEP-1998 (first entry)  
 XX  
 DE Flag-amyloid protein precursor (FLAG-APP-C100) fusion protein.  
 XX  
 KW Amyloid protein precursor; APP-C100; fusion protein; FLAG-APP-C100;  
 KW transgenic animal; Alzheimer's disease; mouse; neurodegeneration;  
 KW brain; ss.  
 XX  
 OS Chimeric - Synthetic.  
 OS Chimeric - Mus sp.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 1..351  
 FT /tag= a  
 FT /product= "FLAG-APP-C100"  
 FT /note= "no start or stop codon specified"  
 XX  
 PN W09816627-A1.  
 XX  
 PD 23-APR-1998.  
 XX  
 PF 16-OCT-1997; 97WO-US18202.  
 XX  
 PR 16-OCT-1996; 96US-0729345.  
 XX  
 PA (MCLE-) MCLEAN HOSPITAL CORP.  
 PA (WELL-) WELLESLEY COLLEGE.  
 XX  
 PI Berger-Sweeney J, Neve RL;  
 XX

DR WPI: 1998-251279/22.  
 DR P-PSDB; AAW51102.  
 XX  
 PT Transgenic animal with rapid development of Alzheimer disease  
 PT characteristics - contains nucleic acid encoding fragment of amyloid  
 PT protein precursor fused to flag peptide, useful as model for  
 PT screening potential therapeutic agents  
 XX  
 PS Claim 5; Pages 26-27; 41pp; English.  
 XX  
 CC This is the nucleotide sequence of the FLAG-Amyloid protein precursor  
 CC (FLAG-APP-C100), used in the method of the invention, involving the  
 CC creation of an FLAG-amyloid protein precursor fusion protein, as well  
 CC as a transgenic animal with rapid development of Alzheimer's disease.  
 CC The animals, specifically mice, show increased neurodegeneration in the  
 CC brain and cognitive impairment, so are used as in vivo models for  
 CC screening agents potentially useful in treatment of Alzheimer's disease.  
 XX  
 SQ Sequence 354 BP; 107 A; 71 C; 95 G; 81 T; 0 other;  
 XX  
 Query Match 22.0%; Score 26.6; DB 19; Length 354;  
 Best Local Similarity 58.0%; Pred. No. 12;  
 Matches 47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;  
 OY 11 TAGCCTCGGTGACAGATCGATGCTGATTAATGAGCGTCCCTGATCCATTG 70  
 DB 176 TAGCGACAGTGTGCTGATCACTTGATGATGATGAAGAAAGATACATCATTC 235  
 OY 71 TTGCTGTGTATTATTAATGTTG 91  
 DB 236 ATCATGTTGTGCTGGAGGTTG 256  
 XX  
 RESULT 45  
 AAV20379  
 ID AAV20379 standard; DNA; 354 BP.  
 XX  
 AC AAV20379;  
 XX  
 DT 26-JUN-1998 (first entry)  
 XX  
 DE DNA for APP C-terminal fragment mutant SPACT Thr43Phe.  
 XX  
 KW Human; amyloid precursor protein; APP; carboxy-terminal fragment;  
 KW A4CT; mutant SPACT Thr43Phe; disease model; treatment;  
 KW Alzheimer's disease; ss.  
 XX  
 OS Homo sapiens.  
 OS Synthetic.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 1..354  
 FT /tag= a  
 FT /note= "stop codon not given"  
 XX  
 PN W09803643-A2.  
 XX  
 PD 29-JAN-1998.  
 XX  
 PF 17-JUL-1997; 97WO-EP03960.  
 XX  
 PR 08-MAY-1997; 97GB-0009239.  
 PR 22-JUL-1996; 96GB-0015351.  
 PR 09-SEP-1996; 96GB-0018804.  
 XX  
 PA (SMK ) SMITHKLINE BEECHAM AUSTRALIA PTY LTD.  
 PA (SMK ) SMITHKLINE BEECHAM PHARMA GMDH.  
 XX  
 PI Beyreuther K, Lichtenhaer S, Masters CL, Prior P;  
 XX  
 DR WPI: 1998-120768/11.  
 DR P-PSDB; AAW50029.  
 XX

Construct containing sequence for mutant form of amyloid precursor protein - or its C-terminal fragment, and related transgenic animals or transformed cells, used for identifying potential drugs for Alzheimer's disease

Claim 10; Page 10; 15pp; English.

The present sequence encodes the human amyloid precursor protein (APP) C-terminal fragment (A4CT) mutant SPACr Thr43Phe. The mutation results in a higher ratio of beta A4 1-40, useful in disease models to identify potential drugs for the treatment of Alzheimer's disease.

Sequence 354 BP; 89 A; 83 C; 99 G; 83 T; 0 other;

Query Match 22.0%; Score 26.6; DB 19; Length 354;

Best Local Similarity 58.0%; Pred. No. 12; Matches 47; Conservative 0; Mismatches 34; Indels 0; Gaps 0;

Qy 11 TAGCCTGGGTGGACAGATCGAGCGCTGCCCATGCTGATAAATGGAGCGGCTGATCCATTG 70  
 ||| | ||| | || | ||| || | || | ||| |||  
 Db 179 TAGCATTCCTGATCCGATCATTGCTGTGATGCTGAGAGAGAAACAGTACACATTCATTG 238

Qy 71 TTCGTTGTGTATTAATGTTG 91  
 || | ||| | | |||  
 Db 239 ATCATGTGTGTGGGAGGTTG 259

Search completed: April 23, 2003, 15:02:20  
 Job time : 127.069 secs

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related to suppress  
hypothetical prote

hypothetical prote

eyelid - fruit fly  
probable drug resi  
glycolate oxidase

carboxylase - Dehn  
hypothetical prote  
chromosome condens  
probably polypeptid

hypothetical prote  
phosphatidate cyti  
hypothetical prote  
transcription fact  
hypothetical prote  
100 kDa coactivato

polymorphic membra  
polymorphic outer  
polymorphic outer  
burnt-bat-fal] are to

hypothetical prote  
conserved hypothet  
related to water

dihydroorotate oxidase  
corticotropin-releasing hormone  
corticotropin-releasing hormone  
corticotropin-releasing hormone

rain PCC 7120)

to, S.; Watanabe, A.  
ada, M.; Yasuda, M.  
change 30-Jun-2002

oxygen-fixing Cyanobacteria

32337; GSPDB:GN0017  
CTGATCCATTGT 71  
:::

Db 412 SerLeuGlyTyrGlnTyrAspAsnProAsnSerIysSerTirpleuGlnPInhevalArgAla 433  
QY 72 TCCTGTCGTATATATGTCATATATGACGACGACACACACCTAG 119  
:::||||| ||| |||||||||:::||||::: ||:::  
Db 432 AsnLeuTyrTyrGlnAsnAlaIleIleGluGluAspSerIAsnArgSer 447

RESULT 2  
1180797

hypothetical protein SA0236 [Imported] - *Staphylococcus aureus* (strain N315)  
C:Species: *Staphylococcus aureus*  
C:Date: 10-May-2001 #sequence\_revision 10-May-2001 #text\_change 22-Oct-2001  
C:Accession: H89787  
R:Kiroda, M.; Ohta, T.; Uchiyama, I.; Baba, T.; Yuzawa, H.; Kobayashi, I.; Cui, L.; Oguc, A.; Mitani-Uli, Y.; Kobayashi, N.; Sawano, T.; Inoue, R.; Kaito, C.; Sekimizu, K.;  
C.: Shiba, T.; Hattori, M.; Ogasawara, N.; Hayashi, H.; Hiramatsu, K.  
A:Title: Whole genome sequencing of methicillin-resistant *Staphylococcus aureus*.  
A:Reference number: A89758; MUID:21311952; PMID:11418146  
A:Accession: H89787  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 17155 <RUR>  
A:Cross-references: GB:BA000018; FID:913700160; PIDN:BA041459.1; GSPDB:GN00149  
A:Experimental source: strain N315  
C:Genetics:  
C:Gene: SA0236

Alignment Scores:

Pred. No.:	3,65	Length:	155
Score:	58.50	Matches:	14
Percent Similarity:	57.14%	Conservative:	10
Best Local Similarity:	33.33%	Mismatches:	11
Query Match:	27.08%	Indels:	7
DB:	2	Gaps:	2

US-09-198-779B-1\_COPY\_1\_121 (1-121) x H89787 (1-155)

0Y 113 TGTGTGTCCTGCTCAATTATACACATTA-----ATAACACAACGAACA----- 69

Db 61 CysSerValAlaIleProHisThrAspValGluHisIleAsnHisArgThrIleGlyVal 80

68 -----ATGGATCAGGACCGTCCATTATCAGCATGGGACGCTGATCTGTCCACCGAG 15

01 ALAVALLEVALYSGUVALPIOPHIELLEGLUMELGYINLEUASPGINGININRGLU 100

1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100
---	---	---	---	---	---	---	---	---	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	----	-----

### RESULT 3

ribosomal protein S2-related protein - *Deinococcus radiodurans* (strain R1)  
C:Species: *Deinococcus radiodurans*  
C:Date: 03-Dec-1999 #sequence-revision 03-Dec-1999 #text-change 28-Jul-2000  
C:Accession: G75298  
R:White, O.; Eisen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.; M.; Shen, M.; Vamathavan, J.J.; Lam, P.; McDonald, L.; Utlacker, T.; Zalewski, C.; Smith, H.O.; Venter, J.C.; Fraser, C.M.  
A:Title: Genome sequence of the radioresistant bacterium *Deinococcus radiodurans* R1.  
Reference number: A75250; MUID:20036896; PMID:10567266

A: Accession: G75298  
A: Status: preliminary

A: Molecule type: DNA  
A: Residues: 1-472 <WH

A: Experimental source:  
A;C:cross-references: GB;  
A:Experimental source:

C;GeneLCS:  
A:Gene: DR2241

C:Superfamily: Deinococcus radiodurans hypothetical protein DBA0012

**Alignment Scores:**

Pred. No.:	4.41	Length:	472
Score:	58.00	Matches:	10
Percent Similarity:	64.71%	Conservative:	1
Best Local Similarity:	58.82%	Mismatches:	6
Query Match:	26.98%	Indels:	0
DB:	2	Gaps:	0

US-09-198-779B-1\_COPY\_1\_121 (1-121) x G75298 (1-472,

18 GGTGGACAGATCGACGCTGCCCATGCTGATAATGGACGGTCCGTATCCAT 68

Db 268 G1YG1YAsPRLeAsPrgInAlaH1SA1aG1ua1aT1rPa1aAlaLeuArgH1s 284

RESUL  
T17243

hypothetical protein DKFZ05860217.1 - human (fragment)  
C:Species: Homo sapiens (man)  
C:Date: 15-Oct-1999 #sequence\_revision 15-Oct-1999 #text\_change 15-Oct-1999  
C:Accession: F17243  
R:Koehnert, K.; Beyer, A.; Mewes, H.W.; Gassenhuber, J.; Wiemann, S.  
submitted to the Protein Sequence Database, September 1999  
A:Reference number: Z18722  
A:Accession: F17243  
A:Status: preliminary  
A:Molecule type: mRNA  
A:Residues: 1-701<KOE>

A; Experimental source:

C:Genetics:		
A:Note: DKFZP586B0217.1		
Alignment Scores:		
Pred. No.:	8.87	Length:
Score:	56.00	Matches:
		14

Best Local Similarity:	36.84%	Mismatches:	10
------------------------	--------	-------------	----

DB:	2	Gaps:	1
-----	---	-------	---

US-09-198-779B-1\_COPY\_1\_121 (1-121) X T17243 (1-701)

QY 104 CTGCTCAATTATACACATTAATAACACAACGAACAATGGATCAGGACCGT----- 54

Db 302 LeuLysAsnPhenylTyrThrLeuValThrGlnArgThrLeuAspArgGluSerGlnAlaGlu 321

CCATTATCAGCATGGGACGCTCGATCTGTCCACCGAG 15  
|||||::|  
23

## RESULT 5

hypothetical protein C05D9.4 - *Caenorhabditis elegans*  
 C.Species: *Caenorhabditis elegans*  
 C.Date: 22-Oct-1999 #sequence\_revision 22-Oct-1999 #text\_change 29-Oct-1999  
 C.Accession: T31059  
 R.Fulton, L. Gartung, S.  
 submitted to the EMBL Data Library, September 1999  
 .Description: The sequence of C. elegans cosmid C05D9.

A;Reference number: Z20960

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Residues: 1-86 <FUL>  
A;Cross-references: EMBL:U64605. PIDN:AA004585 1

A; Experimental source: strain Bristol NZ

A;map position: x  
A;Introns: 19/1: 53/3

A;NOLe: C03D9.4

Alignment Scores:

Pred. No.:	12.2	Length:	86
Score:	55.00	Matches:	12











C:Genetics:1603  
A:Gene: MTH1603  
A:Start codon: TAC  
C:Superfamily: acetate-CoA ligase homology  
C:141-400/Domain: acetate-CoA ligase homology  
#status atypical <ACL>

Alignment Scores:

Pred. No.:	24.9	Length:	400
Score:	53.00	Matches:	9
Percent Similarity:	66.67%	Conservative:	3
Best local Similarity:	50.00%	Mismatches:	6
Query Match:	24.65%	Indels:	0
DB:	2	Gaps:	0

US-09-198-779B-1\_COPY\_1\_121 (1-121) x B69081 (1-400)

```

Oy      26  GATGCACCGCTGCCCATGCTGATAAATGGACGGCTCTGATCCATTGTTCCGTTGTG  79
        |||      |||||||      |||      |||||||      |||      |||
Db      256  ASPASpGLucyspProcysgInGLuLeuAspProGLuAspProLeuPheIleLeu  273

```

RESULT 20

hypothetical protein T12G3.2 - *Caenorhabditis elegans*

C:Protein: AminoAcidSequence\_Cys368  
C>Date: 15-Oct-1999 #sequence\_revision 15-Oct-1999 #text\_change 15-Oct-1999  
C:Accession: T24868  
R:Stims, M.  
submitted to the EMBL Data Library, January 1996  
A:Reference number: Z19945

**Alignment Scores:**

Pred. No.:	25.2	length:	922
Score:	53.00	Matches:	14
Percent Similarity:	61.29%	Conservative:	5
Best Local Similarity:	45.16%	Mismatches:	6
Query Match:	24.54%	Indels:	6
DB:	2	Gaps:	2

US-09-198-779B-1\_COPY\_1\_121 (1-121) x T24868 (1-928,

```

Oy 105 CCTGTCATTTATACACATTAAACACACG-----AACCATGGATCA 61
    |||:::||||| ::| ||| ||||| |||||
Db 895 ProserGlnLeuProserProAsnGlnThrIleProHisMetTyAsnAsnGlyIle 91

```

oy 60 GGACCTCCATTATCAGCATGGGAGCGTCGA 28  
 ||| |||||::||| |||:::  
 Db 915 ---PrometiletyrGluHisHisGlnGln 924

## RESULT 21

hypothetical protein PFB0540w - malaria parasite (Plasmodium falciparum)

C:Species: Plasmodium falciparum  
C:Date: 13-Nov-1998 #sequence\_revision 13-Nov-1998 #text\_change 21-Jul-2000  
C:Accession: D11612  
R:Gardner, M.J., Tetteh, H., Carucci, D.J., Cummings, L.M., Aravind, L., Koonin, E.V.,  
Pierse, M., Salzberg, S., Zhou, L., Sutton, G.G., Clayton, R., White, O., Smith, H.O.  
Science 282, 1126-1133, 1998  
A>Title: Chromosome 2 sequence of the human malaria parasite Plasmodium falciparum.  
A:Reference number: A1600; MUID:99021743; PMID:9804551

A;Cross-references: GB:AE001401; GB:AE001362; MID:g3845209; PIDN:AAC71897.1; PID:g3845209  
A;Experimental source: clone 3D7  
C;Genetics:  
A;Gene: PFB0540w

Alignment Scores:

Pred. No.:	25.4	Length:	184
Score:	53.00	Matches:	10
Percent Similarity:	53.85	Conservative:	4
Best local Similarity:	38.46	Mismatches:	12
Query Match:	24.54	Indels:	0
DB:	2	Gaps:	0

US-09-198-779B-1\_COPY\_1\_121 (1-121) x D71612 (1-1844)

QY 90 AACATTATATACCAACCAACAATGATGATCAGGACCGTCATTATATCAGCATGGGACGGT 31  
 ||| |||| |||| :: ||||  
 Db 881 AsnAsnAsnAsnAsnAsnAsnAsnAsnAsnHisSerIleIleAsnAsnAsnIleThr 90

```
QY      30 CGATCTGTCCACCGAGGC 13
          :: ::| | | | | | |
Db      901 GlnGlyIleHisArgGly 906
```

RESULT 22

eyelid - fruit fly (*Drosophila melanogaster*)

c;Species: *Esopharia melanogaster*;  
 C.Date: 13-Aug-1999 #sequence\_revision 13-Aug-1999 #text\_change 17-Nov-2000  
 C.Accession: T13049  
 R:Teisman, J.E.; Luk, A.; Rubin, G.M.; Heberlehn, U.  
 Submitted to the EMBL Data Library, March 1998  
 A.Reference number: Z17592

## Alignment Scores

Pred. No.:	25	6	Length:	271
Score:	53.00		Matches:	10
Percent Similarity:	78.57%		Conservative:	1
Best Local Similarity:	71.43%		Mismatches:	3
Query Match:	24.54%		Indels:	0
DB:	2		Gaps:	0

US-09-198-779B-1\_COPY\_1\_121 (1-121) x T13049 (1-2715)

```

QY      84 AATAACACACGACGAACATGGATCAGGACCGTCATTATATAG 433
        ||||| :::||||| ||||| ||||| . |||
Db     46 AsnAsnAsnSerAsnAsnGlySerAsnProSerIleGlnGln 50

```

## RESULT 23

probable drug resistance protein [imported] - Sinorhizoc

C:Species: *Sinorhizobium meliloti*  
C:Date: 24-Aug-2001 #sequence\_revision: 24-Aug-2001 #text\_change: 30-Sep-2001  
C:Accession: G95375  
R:Barrett, M.J., Fisher, R.F., Jones, T., Komp, C., Abola, A.P., Barloy-Hubler, F.,  
Kaltman, S., Keeling, D.H., Palm, C., Peck, M.C., Surzycki, R., Wells, D.H., Yeh,  
Proc. Natl. Acad. Sci. U.S.A. 98, 9883-9888, 2001  
A:Title: Nucleotide sequence and predicted functions of the entire *Sinorhizobium meli*  
A:Reference number: A95262; MUID: 21396509; PMD: 11481432

A:Cross-references: GB:AE006469; PIDN:BAK5569.1; PID:G14524048; GSPDB:GN00165  
A:Experimental source: strain 1021, megaplasmid pSymA  
R:Galibert, F.; Filan, T.M.; Long, S.R.; Puhler, A.; Abola, P.; Ampe, F.; Bailly-Huiler  
pela, D.; Chain, P.; Cowie, A.; Davis, R.W.; Dreano, S.; Federspiel, N.A.; Fisher, R.F.;  
L.; Hyman, R.W.; Jones, T.  
Science 293, 668-672, 2001  
A:Authors: Kahn, D.; Kahn, M.L.; Kalman, S.; Keating, D.H.; Kiss, E.; Komp, C.; Laure  
habault, P.; Vandenbol, M.; Vorholter, F.J.; Weiner, S.; Wells, D.H.; Wong, K.; Yeh, K  
A:Title: The complete genome of the legume symbiont *Sinorhizobium meliloti*.  
A:Reference number: AB6039; MUID:21368234; PMID:11474104  
A:Contents: annotation  
C:Genetics:  
A:Gene: Sma1664  
A:Genome: plasmid

Alignment Scores:  
Pred. No.: 35.2 Length: 388  
Score: 52.00 Matches: 12  
Percent Similarity: 58.33% Conservative: 9  
Best Local Similarity: 33.33% Mismatches: 14  
Query Match: 24.07% Indels: 1  
DB: 2 Gaps: 0

US-09-198-779B-1\_COPY\_1\_121 (1-121) x G95375 (1-388)

OY 114 GTGTTGTGCTGCTCAATTATACACATTATTAACACACAGACAGATGATCAGGACCG 55  
:::||||| ||| ||| ::| ||::||| ::|  
DB 278 LeuLeuMePrGjYgLn-PheValThrValIleValGIuProGIuGIuArgGIuGIuAr 297  
:::||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

OY 54 TCCATTATACAGACGCGTCGATGTCGCCACCGACGCGTAA 9  
||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

DB 297 gProValValProValGIuSerValGIuSerValGIuInaSPargGIuGIuArg 312

RESULT 24  
G83916  
glycolate oxidase iron-sulfur subunit BH2135 [imported] - Bacillus halodurans (strain C-  
C:Species: Bacillus halodurans  
C:Date: 01-Dec-2000 #sequence\_revision 01-Dec-2000 #text\_change 15-Jun-2001  
C:Accession: G83916  
R:Takami, H.; Nakasone, K.; Takaki, Y.; Maeno, G.; Sasaki, R.; Masui, N.; Fujii, F.; Hira  
no, T.  
Nucleic Acids Res. 28, 4317-4331, 2000  
A:Title: Complete genome sequence of the alkaliphilic bacterium Bacillus halodurans and  
A:Reference number: AB36550; MUID:20512582; PMID:11058132  
A:Accession: G83916  
A:Status: preliminary  
A:Molecule type: DNA  
A:Residues: 1-451 <STO>  
A:Cross-references: GB:AP001514; GB:BA000004; NID:q10174613; PIDN:BA005854.1; GSPDB:GN00  
C:Genetics:  
A:Experimental source: strain C-125  
A:Gene: BH2135  
C:Superfamily: Synecchocystis (S)-2-hydroxy-acid oxidase

Alignment Scores:  
Pred. No.: 35.2 Length: 451  
Score: 52.00 Matches: 11  
Percent Similarity: 45.16% Conservative: 3  
Best Local Similarity: 35.48% Mismatches: 15  
Query Match: 24.19% Indels: 2  
DB: 2 Gaps: 1

US-09-198-779B-1\_COPY\_1\_121 (1-121) x G83916 (1-451)

OY 6 CCGTTACCGTCGGTGAGACAGATCGACCGCTGCCCATGCTGAT-----AAATGACGCGTC 59  
||||| ||| ||| ::| ||| |||  
DB 11 ProProSeIleThrAsnTyrHisTTrpSerAspHisProAspProAsnLysTrpAlaasp 30  
:::||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

OY 60 CTGATCCATTGCTGCTGCTGTTATTAATGTTGT 92  
:::||||| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::| ::|

DB 31 CysValHisCysGlyMetCysLeuGluAlaCys 41

RESULT 25  
S17998

gene COX1 intron 4 protein - Yeast (Kluyveromyces marxianus var. lactis) mitochondrial D  
C.Species: mitochondrion Kluyveromyces marxianus var. lactis, Candida sphaerica  
C.Date: 10-Mar-1994 #sequence\_revision 10-Mar-1994 #text\_change 08-Sep-2000  
R.Hardy, C.M.; Clark-Walker, G.D.  
Curr. Genet. 20, 99-114, 1991  
A.Title: Nucleotide sequence of the COX1 gene in Kluyveromyces lactis mitochondrial D  
A.Reference number: S17993; MUID:92035081; PMID:1657415  
A.Accession: S17998  
A.Molecule type: DNA  
A.Residues: 1-763 <HAR>  
A.Cross-references: EMBL:X57546  
C.Genetics:  
A.Genome: mitochondrion  
A.Genetic code: SGC2  
A.Introns: 69/1, 235/3, 324/2  
C.Superfamily: cytochrome c oxidase chain I homology  
C.Keywords: mitochondrion  
F:1-369/Region: cox1 exons 1 to 4 encoded  
F:370-763/Region: cox1 intron encoded

Alignment Scores:  
Pred. No.: 35.5 Length: 763  
Score: 52.00 Matches: 9  
Percent Similarity: 75.00% Conservative: 3  
Best Local Similarity: 56.25% Mismatches: 4  
Query Match: 24.07% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1\_COPY\_1\_121 (1-121) x S17998 (1-763)

QY 114 GGTGTTGTCTGCTCAATTATACACATTATACACACGACCAAT 67  
:::||||| :|||||  
Db 408 ILeuTYrSerIleYsMetIlyrAsnIleAsnAsnAsnAsn 423

RESULT 26  
C75585  
carboxylase - Deinococcus radiodurans (strain R1)  
C.Species: Deinococcus radiodurans  
C.Date: 03-Dec-1999 #sequence\_revision 03-Dec-1999 #text\_change 17-Mar-2000  
C.Accession: C75585  
R.White, O.; Eissen, J.A.; Heidelberg, J.F.; Hickey, E.K.; Peterson, J.D.; Dodson, R.J.  
M.; Shen, M.; Vamathevan, J.J.; Lam, P.; McDonald, L.; Utterback, T.; Zaleski, C.;  
S.; Smith, H.O.; Venter, J.C.; Fraser, C.M.  
Science 286, 1571-1577, 1999  
A.Title: Genome sequence of the radioreistant bacterium Deinococcus radiodurans R1.  
A.Reference number: A75250; MUID:20036896; PMID:10567266  
A.Accession: C75585  
A.Status: preliminary  
A.Molecule type: DNA  
A.Residues: 1-1091 <WHI>  
A.Cross-references: GB:AE001863; GB:AE001825; NID:96460670; PIDN:AAF12468.1; PID:9646  
A.Experimental source: strain R1  
C.Genetics:  
A.Gene: DRA0310  
A.Map position: 2

Alignment Scores:  
Pred. No.: 35.7 Length: 1091  
Score: 52.00 Matches: 11  
Percent Similarity: 80.00% Conservative: 1  
Best Local Similarity: 73.33% Mismatches: 3  
Query Match: 24.19% Indels: 0  
DB: 2 Gaps: 0

US-09-198-779B-1\_COPY\_1\_121 (1-121) x C75585 (1-1091)

QY 6 CCGTCTAGCCTCGGTGAGACAGATGACGCTGCCCATGTGATTA 50  
||||| ||||||| ||||||| |||||||  
Db 902 ProAlaValleuGLyGLyAlaIleAspAlaIleagIyAlaAspIys 916

RESULT 27  
548424

```

Hypothetical protein YII055c - Yeast (Saccharomyces cerevisiae)
C:Species: Saccharomyces cerevisiae
C:Date: 02-Dec-1994 #sequence_revision 02-Dec-1994 #text_change 19-Apr-2002
C:Accession: S48424
R:Smith, V.
submitted to the EMBL Data Library, September 1994
A:Reference number: S48407
A:Accession: S48424
A:Molecule type: DNA
A:Residues: 1-627 <SMI>
A:Cross-references: GB:247047, EMBL:Z38060, NID:g603997, PID:g763291, GSPDB:GN00009, MIF
C:Genetics:
A:Gene: MIPS:YII055c
A:Cross-references: SGD:S0001317
A:Map position: 9L

Alignment Scores:
Pred. No.: 42.1 Length: 627
Score: 51.50 Matches: 11
Percent Similarity: 64.29% Conservative: 7
Best Local Similarity: 39.29% Mismatches: 7
Query Match: 23.84% Indels: 3
DB: 2 Gaps: 1

US-09-198-7798-1_COPY_1_121 (1-121) x S48424 (1-627)
OY 96 TTATACACATTATAT-----AACACAGCAACATGATCAGACCGTCATTAT 46
|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 329 LeuYrAnleuAsnMeTAlSerAsnSerAsnAsnAngLYAsnIleProThrThnSer 348
OY 45 CAGCATGGCGAGACGTCGATCTGTC 22
::|||:|||||:
Db 349 ThrAnGlyAspAspArgAlaLeu 356

RESULT 28
T24216
chromosome condensation protein homolog DPY-27 - Caenorhabditis elegans
N:Alternate names: protein R13G10.1
C:Species: Caenorhabditis elegans
C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 02-Jun-2000
C:Accession: T24216; A55095
R:Gardner, A.
submitted to the EMBL Data Library, August 1994
A:Reference number: Z19857
A:Accession: T24216
A:Status: translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-1469 <MIL>
A:Cross-references: EMBL:Z35602; PIDN:CAA84669.1; GSPDB:GN00021; CESP:R13G10.1
A:Experimental source: clone R13G10
R:Chuang, P.T.; Albertson, D.G.; Meyer, B.J.
Cell 79, 459-474, 1994
A:Title: Dpy-27: a chromosome condensation protein homolog that regulates Caenorhabditis
A:Reference number: A55095; MUID:95042743; PMID:7954812
A:Accession: A55095
A:Status: preliminary; nucleic acid sequence not shown
A:Molecule type: DNA
A:Residues: 1-1132/'G', 1134-1469 <CHU>
A:Cross-references: GB:IJ5274; NID:g529384; PIDN:AAA62647.1; PID:g529385
A:Note: authors translated the codon GAT for residue 1133 as Gly
C:Genetics:
A:Gene: CESP:R13G10.1
A:Map position: 3
A:Introns: 142/3; 176/3; 296/2; 548/2; 627/3; 697/2; 847/3; 993/3; 1086/2; 1202/3; 1383/3;
C:Superfamily: chromosome segregation protein SMC1

Alignment Scores:
Pred. No.: 42.6 Length: 1469
Score: 51.50 Matches: 12
Percent Similarity: 57.69% Conservative: 3
Best Local Similarity: 46.15% Mismatches: 8
Query Match: 23.84% Indels: 3
DB: 2 Gaps: 1

```

```
US-09-198-779B-1_COPY_1_L121 (1-121) x Tt24216 (1-1469)
```

```
OY      71 ACATGTCACAGCACCCTCCATTATC-----AGCATGGCGACGCATGTCTGCC    21  
          ||| ||| ||||||||:::  
Db       10 ThSeraSPasPArGPProTYrAlAsPhnRASPserMetPrGIUValASpleuAasp   29  
  
OY      20 ACCGAGGCTACAGCGAAA     3  
          ::: ||||||::::  
Db       30 ValASpaTGALgARtgIn 35
```

RESULT 29

E70522

probable polyketide synthase - Mycobacterium tuberculosis (strain H37RV)  
C:Species: Mycobacterium tuberculosis  
C>Date: 17-Jul-1998 #sequence\_revision 17-Jun-1998 #text\_change 20-Jun-2000  
C:Accession: E70522

R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon,  
J.; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd,  
Raftandram, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skellton, S.; Squares, S.  
Nature 393, 537-544, 1998

A:Authors: Squares, R., Sulston, J.E., Taylor, K., Whitehead, S., Barrell, B.G.

A>Title: Deciphering the biology of Mycobacterium tuberculosis from the complete genome sequence  
A:Reference number: A70500; PMID:98295987; PMID:9634230

A:Accession: E70522

A>Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-2126 <COL>

A:Cross-references: GB:297188; GB:A123456; NID:g3z61805; PIDN:CAB10012.1, PID:g22248

A:Experimental source: Strain H37RV

C:Genetics:

A:Gene: pksZ

C:Superfamily: mycoesteroid acid synthase: 3-oxoacyl-[acyl-carrier-protein] synthase I  
name homology; [acyl-carrier-protein] S-malonyltransferase homology

C:Keywords: carrier protein

F:46-445/Domain: 3-oxoacyl-[acyl-carrier-protein] synthase I homology <OAS>  
F:553-833/Domain: [acyl-carrier-protein] S-malonyltriferase homology <AMT>  
F:1451-1739/Domain: long-chain alcohol dehydrogenase homology <ADH>  
F:1772-1952/Domain: short-chain alcohol dehydrogenase homology <SDH>  
F:2044-2109/Domain: acyl carrier protein homology <ACP>

C:Genetics:  
 A:Map position: 4  
 A:Introns: 69/1; 83/3; 123/3; 176/2; 201/2; 224/3; 275/3; 312/1; 333/1  
 A>Note: T16L1.30  
 C:Superfamily: Arabidopsis thaliana hypothetical protein T16L1.30

## Alignment Scores:

Pred. No.:	Length:	Matches:	Conservative:
Score:	49.7	371	12
Percent Similarity:	51.00	60.87%	2
Best Local Similarity:	52.17%	Mismatches:	9
Query Match:	23.72%	Indels:	0
DB:	2	Gaps:	0

US-09-198-779b-1\_COPY\_1\_121 (1-121) x T04971 (1-371)

OY 19 GTGACAGATGACGCTGCTGCTGATTAATGAGCGCTCCTGATTCATTGTCGTTGT 78

Db 153 ValAspIysAspThrIleuProValIeuPheAsnProIeuSerPheHisPhePheArgMet 172

OY 79 GTTATTAAAT 87

Db 173 ValIleAsn 175

RESULT 31

T47873 phosphatidate cytidyltransferase-like protein - Arabidopsis thaliana

N:Alternate names: protein T4C21.30

C:Species: Arabidopsis thaliana (mouse-ear cress)

C>Date: 20-Apr-2000 #sequence\_revision 20-Apr-2000 #text\_change 20-Apr-2000

C:Accession: T47873

R:Choisne, N.; Robert, C.; Brottier, P.; Wincker, P.; Catolico, L.; Artiguenave, F.; S.

W.; Rudd, S.; Lemcke, K.; Mayer, K.F.X.

Submitted to the Protein Sequence Database, March 2000

A:Reference number: 224479

A:Accession: T47873

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-399 <CHO>

A:Cross-references: EMBL:AL162295

A:Experimental source: cultivar Columbia; BAC clone T4C21

C:Genetics:

A:Map position: 3

A:Introns: 81/3; 185/2; 249/1; 290/3; 335/2; 374/1

A>Note: T4C21.30

Alignment Scores:

Pred. No.:	Length:	Matches:	Conservative:
Score:	49.7	399	16
Percent Similarity:	51.00	52.78%	3
Best Local Similarity:	44.44%	Mismatches:	11
Query Match:	23.72%	Indels:	6
DB:	2	Gaps:	2

US-09-198-779b-1\_COPY\_1\_121 (1-121) x T47873 (1-399)

OY 3 TTTCGCTTACCTCGGCGACATGACCTGCTGATTAATGACGCTCTG 62

Db 254 TTPProThrIleLeuGlyGln-----AlaHis-----TTPThrValGly 267

OY 63 ATCCATTGTCGTTGTTATTATTATGTTTAAATGAGACAGACACA 110

Db 268 LeuValAlaIleLeuIleSerPheCysGlyIleIleAlaSerAspThr 283

RESULT 32

T32605 hypothetical protein E04A4.3 - Caenorhabditis elegans

C:Species: Caenorhabditis elegans

C>Date: 29-Oct-1999 #sequence\_revision 29-Oct-1999 #text\_change 29-Oct-1999

C:Accession: T32605

R:Sammons, L.; Wohldmann, P.; Biewald, T.

submitted to the EMBL Data Library, December 1997

A:Description: The sequence of C. elegans cosmid E04A4.

A:Reference number: 221199

A:Accession: T32605

A:Status: preliminary; translated from GR/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-433 <SAM>

A:Cross-references: EMBL:AF038611; PIDN:AAB92038.1; GSFDB:GN00022; CESP:E04A4.3

A:Experimental source: strain Bristol N2; clone E04A4

C:Genetics:

A:Gene: CESP:E04A4.3

A:Map position: 4

A:Introns: 21/2; 113/1; 147/3; 190/1; 283/2; 345/3; 382/3; 418/2

Alignment Scores:

Pred. No.:	Length:	Matches:	Conservative:
Score:	49.8	433	10
Percent Similarity:	51.00	57.89%	1
Best Local Similarity:	52.63%	Mismatches:	8
Query Match:	23.72%	Indels:	0
DB:	2	Gaps:	0

US-09-198-779b-1\_COPY\_1\_121 (1-121) x T32605 (1-433)

OY 20 TGGACAGATGACGCTGCTGCTGATTAATGAGCGCTCCTGATTCATTGTCGTT 76

Db 324 TTPGlyGlyArgArgCysProCysLeuValAspGlyIuaAlaIleVal 342

RESULT 33

JQ2010 transcription factor POU-1 - planarian (Dugesia japonica)

C:Species: Dugesia japonica

C>Date: 03-Feb-1994 #sequence\_revision 03-Feb-1994 #text\_change 16-Jun-2000

C:Accession: JQ2010

R:Orli, H.; Agata, K.; Watanabe, K.

Biochem. Biophys. Res. Commun. 192, 1395-1402, 1993

A:Title: POU-domain genes in planarian Dugesia japonica

A:Reference number: JQ2010; MUID:93282851; PMID:8099480

A:Accession: JQ2010

A:Molecule type: mRNA

A:Residues: 1-559 <ORI>

A:Cross-references: DDBJ:DJ12924; NID:9217311; PIDN:BAA02308.1; PID:9217312

A:Experimental source: strain GI

C:Genetics:

A:Gene: DpOU1

C:Superfamily: planarian transcription factor POU-1; homeobox homology; POU domain ho

C:Keywords: DNA binding; homeobox; nucleus; transcription regulation

F:266-333/Domain: POU domain homology <POU>

F:352-408/Domain: homeobox homology <HOX>

Alignment Scores:

Pred. No.:	Length:	Matches:	Conservative:
Score:	50	559	8
Percent Similarity:	51.00	58.33%	6
Best Local Similarity:	33.33%	Mismatches:	10
Query Match:	23.61%	Indels:	0
DB:	1	Gaps:	0

US-09-198-779b-1\_COPY\_1\_121 (1-121) x JQ2010 (1-559)

OY 105 CCTGCTCAATTATACACATTATACACAGCAATGATGACGCTCCATTAT 46

Db 534 ProArgSerPhePheMetIleYasAsnIleYerIersnGlyThrThrProThrLeuPhe 553

OY 45 CAGCATGGCAG 34

Db 554 AspHisSerGln 557

RESULT 34

T18649 hypothetical protein B0024.14 - Caenorhabditis elegans

C:Species: Caenorhabditis elegans

C>Date: 15-Oct-1999 #sequence\_revision 15-Oct-1999 #text\_change 15-Oct-1999

C:Accession: T18649

R:McMurray, A.

submitted to the EMBL Data Library, April 1996

A:Reference number: Z19001

A:Accession: T18649

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-884 <WILL>

A:Cross-references: EMBL:Z11178; PIDN:CAA94886.1; GSPDB:GN00023; CESP:B0024.14

A:Experimental source: clone B0024

C:Genetics:

A:Gene: CESP:B0024.14

A:Map position: 5

A:Introns: 46/3; 84/1; 212/1; 307/2; 345/2; 394/1; 424/2; 481/1; 596/1; 702/1; 765/3; 85

Alignment Scores:

Pred. No.:	50.3	Length:	884
Score:	51.00	Matches:	12
Percent Similarity:	54.84%	Conservative:	5
Best Local Similarity:	38.71%	Mismatches:	14
Query Match:	23.72%	Indels:	0
DB:	2	Gaps:	0

US-09-198-779B-1\_COPY\_1\_121 (1-121) x T18649 (1-884)

OY 3 TTTCGCTACCTCGTGACAGATGCGCCATGCTGATTAATGACGGCTCTG 62

DB 717 PheProThrSerLeuGlyProArgValAspAspGlyAsnGlySerSerAlaThrSerLeu 736

OY 63 ATCCATTCGTGCTGTATTAATGCTGTATA 95

DB 737 IleValValSerLeuMetSerLeuGlyValVal 747

RESULT 35

138968

100 kDa coactivator - human

C:Species: Homo sapiens (man)

C:Date: 29-May-1998 #sequence\_revision 29-May-1998 #text\_change 28-Jul-2000

C:Accession: I38968

R.Tong, X.; Drapkin, R.; Yalamanchili, R.; Mostafaei, G.; Kieff, E.

Mol. Cell. Biol. 15, 4735-4744, 1995

A:Title: The Epstein-Barr virus nuclear protein 2 acidic domain forms a complex with a

A:Reference number: I38968; MUID:95379816; PMID:7651391

A:Accession: I38968

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: mRNA

A:Residues: 1-885 <RES>

A:Cross-references: EMBL:U23055; NID:9799176; PIDN:AA80488.1; PID:9799177

C:Superfamily: Schizosaccharomyces pombe probable transcription factor SPCC645.08c

Alignment Scores:

Pred. No.:	50.3	Length:	885
Score:	51.00	Matches:	11
Percent Similarity:	75.00%	Conservative:	4
Best Local Similarity:	55.00%	Mismatches:	1
Query Match:	23.61%	Indels:	4
DB:	2	Gaps:	1

US-09-198-779B-1\_COPY\_1\_121 (1-121) x I38968 (1-885)

OY 61 AGGACCGTCATTTATCAGCATGGCAGCGTCGATCTGTCACCGACGCTAGACGGA 2

DB 341 LysThrIleHisLeuSer-----SerIleArgProProArgLeuGluGlyGlu 356

RESULT 36

F81539

polymorphic membrane protein G family CP0770 [imported] - Chlamydia pneumoniae (str

C:Species: Chlamydia pneumoniae; Chlamydia pneumoniae

C:Date: 31-Mar-2000 #sequence\_revision 31-Mar-2000 #text\_change 15-Jun-2001

C:Accession: F81539

R.Reed, T.D.; Brunham, R.C.; Shen, C.; Gill, S.R.; Heidelberg, J.F.; White, O.; Hickey,

C.; Dodson, R.; Gwinn, M.; Nelson, W.; Deboy, R.; Kolonay, J.; McClarty, G.; Salzberg,

Nucleic Acids Res. 28, 1397-1406, 2000

A:Title: Genome sequences of Chlamydia trachomatis Morn and Chlamydia pneumoniae AR39.

A:Reference number: AB1500; MUID:20150255; PMID:10684935

A:Accession: F81539

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-922 <REA>

A:Cross-references: GB:AE002237; GB:AE002161; NID:97189684; PIDN:AAF38570.1; PID:9718

A:Experimental source: strain AR39, HL cells

C:Genetics:

A:Gene: CP0770

C:Superfamily: Chlamydia pneumoniae polymorphic outer membrane protein G

Alignment Scores:

Pred. No.:	50.3	Length:	922
Score:	51.00	Matches:	12
Percent Similarity:	58.06%	Conservative:	6
Best Local Similarity:	38.71%	Mismatches:	12
Query Match:	23.61%	Indels:	1
DB:	2	Gaps:	0

US-09-198-779B-1\_COPY\_1\_121 (1-121) x F81539 (1-922)

OY 102 GCTCAATTATACACATTATATACACAGCAATGATGAGACCGTCATTATACG 43

DB 209 AlaGluIleArgPheAlaGlnAsnThrAlaLysAsnGlySerGlyAlaLeuTyr-Se 228

OY 42 CATGGCAGCGTCGATCTGTCACCGAGGCT 12

DB 228 rAspGlyAspIleAspIleAspGlnAsnAla 238

RESULT 37

B72131

polymorphic outer membrane protein g family - Chlamydia pneumoniae (strain CWL029

C:Species: Chlamydia pneumoniae; Chlamydia pneumoniae

C:Date: 23-Apr-1999 #sequence\_revision 23-Apr-1999 #text\_change 15-Jun-2001

C:Accession: B72131

R.Kalman, S.; Mitchell, W.; Marthe, R.; Lammel, C.; Fan, J.; Olinger, L.; Grimwood,

Nature Genet. 21, 385-389, 1999

A:Title: Comparative genomes of Chlamydia pneumoniae and C. trachomatis.

A:Reference number: A72000; MUID:99206606; PMID:10192388

A:Accession: B72131

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-922 <ARN>

A:Cross-references: GB:AE001585; GB:AE001363; NID:94376255; PIDN:AA018163.1; PID:9437

A:Experimental source: strain CWL029

C:Genetics:

A:Gene: pmp.1

C:Superfamily: Chlamydia pneumoniae polymorphic outer membrane protein G

Alignment Scores:

Pred. No.:	50.3	Length:	922
Score:	51.00	Matches:	12
Percent Similarity:	58.06%	Conservative:	6
Best Local Similarity:	38.71%	Mismatches:	12
Query Match:	23.61%	Indels:	1
DB:	2	Gaps:	0

US-09-198-779B-1\_COPY\_1\_121 (1-121) x B72131 (1-922)

OY 102 GCTCAATTATACACATTATATACACAGCAATGATGAGACCGTCATTATACG 43

DB 209 AlaGluIleArgPheAlaGlnAsnThrAlaLysAsnGlySerGlyAlaLeuTyr-Se 228

OY 42 CATGGCAGCGTCGATCTGTCACCGAGGCT 12

DB 228 rAspGlyAspIleAspIleAspGlnAsnAla 238

RESULT 38

B66491

polymorphic outer membrane protein G family [imported] - Chlamydia pneumoniae (st

C:Species: Chlamydia pneumoniae; Chlamydia pneumoniae

C:Date: 02-Mar-2001 #sequence\_revision 02-Mar-2001 #text\_change 15-Jun-2001

C:Accession: B66491

R.Shirai, M.; Hirakawa, H.; Kimoto, M.; Tabuchi, M.; Kishi, F.; Ouchi, K.; Shiba, T.;







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